

QY	61	NSVLNTLIMISIVMVTIFLVLYKRYCYKFINGWLINSLMLFLTYIYLGEVLKTYN	120	181 GAIISYDVLAVYLCPKGPLRMLVETAEQRNEIFPALLYSSAMWTVGMAKLDPSSQGLQ	240
QY	137	NSVLNTLIMISIVMVTIFLVLYKRYCYKFINGWLINSLMLFLTYIYLGEVLKTYN	196	257 GAISYDVLAVYLCPKGPLRMLVETAEQRNEIFPALLYSSAMWTVGMAKLDPSSQGLQ	316
QY	121	VMDYPTLLTWNFGAVGMYC1HWKGP1YVLOQAYLIMISALMAYF1KYLPEWSAWYL	180	241 LPYDPMEEDSYDSFGEPSYFEVFPPLTGYPGEELBEFEERGVKLGLGDFIFYPSVLYGK	300
QY	197	VMDYPTLLTWNFGAVGMYC1HWKGP1YVLOQAYLIMISALMAYF1KYLPEWSAWYL	256	257 GAISYDVLAVYLCPKGPLRMLVETAEQRNEIFPALLYSSAMWTVGMAKLDPSSQGLQ	316
QY	181	GATSVYDVLAVYLCPKGPLRMLVETAEQRNEIFPALLYSSAMWTVGMAKLDPSSQGLQ	240	241 LPYDPMEEDSYDSFGEPSYFEVFPPLTGYPGEELBEFEERGVKLGLGDFIFYPSVLYGK	300
QY	317	LPPDKEEDSYDSFGEPSYFEVFPPLTGYPGEELBEFEERGVKLGLGDFIFYPSVLYGK	376	317 AAATGSGDWNTTLACFVALIIGCLTLLLAVFKKALPALPISITFGLIFYFSTDNLVRP	360
QY	301	AAATGSGDWNTTLACFVALIIGCLTLLLAVFKKALPALPISITFGLIFYFSTDNLVRP	360	301 AAATGSGDWNTTLACFVALIIGCLTLLLAVFKKALPALPISITFGLIFYFSTDNLVRP	360
QY	377	AAATGSGDWNTTLACFVALIIGCLTLLLAVFKKALPALPISITFGLIFYFSTDNLVRP	436	377 AAATGSGDWNTTLACFVALIIGCLTLLLAVFKKALPALPISITFGLIFYFSTDNLVRP	436
QY	361	FMDTFLASHQLYI	372	361 FMDTFLASHQLYI	372
QY	437	FMDTFLASHQLYI	448	437 FMDTFLASHQLYI	448
QY	W28508;	W28508 standard; Protein: 447 AA.	7	W28508;	7
QY	W28508;	07-DEC-1997 (first entry)	W28508;	07-DEC-1997 (first entry)	W28508;
QY	AD3;	Full AD4/AD3LP sequence.	AD3;	Full AD4/AD3LP sequence.	AD3;
QY	Presenilin;	Alzheimer's disease; chromosome: miss segregation; inhibitor: AD; trisomy 21; ss.	Presenilin;	Alzheimer's disease; chromosome: miss segregation; inhibitor: AD; trisomy 21; ss.	Presenilin;
QY	W0970723-A2.	W0970723-A2.	W0970723-A2.	W0970723-A2.	W0970723-A2.
QY	PD	27-FEB-1997.	PD	27-FEB-1997.	PD
QY	PR	15-AUG-1996; U133314.	PR	15-AUG-1996; U133314.	PR
QY	PR	16-AUG-1995; US-002448.	PR	16-AUG-1995; US-002448.	PR
QY	(HARD)	HARVARD COLLEGE.	(HARD)	HARVARD COLLEGE.	(HARD)
QY	LI J.,	Potter H.	LI J.,	Potter H.	LI J.,
QY	WP;	97-165297/15.	WP;	97-165297/15.	WP;
QY	N-PSDB;	T87426.	N-PSDB;	T87426.	N-PSDB;
QY	PT	Identifying genes which cause chromosome miss segregation - useful for identifying causes of and treatments for diseases, e.g. Alzheimer's disease, cancer and ageing	PT	Identifying genes which cause chromosome miss segregation - useful for identifying causes of and treatments for diseases, e.g. Alzheimer's disease, cancer and ageing	PT
QY	PS	Claim 29; Fig 29; 77pp; English.	PS	Claim 4; Page -; 178pp; English.	PS
QY	CC	Screening for inhibitors of chromosome miss segregation and processes caused by genes encoding chromosome miss segregation promoters	CC	Screening for inhibitors of chromosome miss segregation and processes caused by genes encoding chromosome miss segregation promoters	CC
QY	CC	was exemplified using Alzheimer's disease. The sequences given in T87401 to T87426 can be used in the above methods.	CC	was exemplified using Alzheimer's disease. The sequences given in T87401 to T87426 can be used in the above methods.	CC
QY	CC	It is not clear from the figure legend, the figure and the disclosure of the specification which sequence of Fig 1 and Fig 28 is the AD4/AD3LP or the AD3 sequence.	CC	It is not clear from the figure legend, the figure and the disclosure of the specification which sequence of Fig 1 and Fig 28 is the AD4/AD3LP or the AD3 sequence.	CC
QY	SQ	Sequence 447 AA;	SQ	Sequence 447 AA;	SQ
QY	Best Local Similarity	99.2%; Score 1907.5; DB 1; Length 447;	QY	Best Local Similarity	99.2%; Score 1907.5; DB 1; Length 447;
QY	Matches	371; Conservative 0; Mismatches 0; Indels 1; Gaps 1;	QY	Matches	371; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
QY	1	EELTKYGAHKYIMLFYPTVLCMIVVATIKSYRFETYKEKGQLIYTPTEDTPSVGORL	60	1 EELTKYGAHKYIMLFYPTVLCMIVVATIKSYRFETYKEKGQLIYTPTEDTPSVGORL	60
QY	77	EELTKYGAHKYIMLFYPTVLCMIVVATIKSYRFETYKEKGQLIYTPTEDTPSVGORL	136	77 EELTKYGAHKYIMLFYPTVLCMIVVATIKSYRFETYKEKGQLIYTPTEDTPSVGORL	136
QY	61	NSVLNTLIMISIVMVTIFLVLYKRYCYKFINGWLINSLMLFLTYIYLGEVLKTYN	120	61 NSVLNTLIMISIVMVTIFLVLYKRYCYKFINGWLINSLMLFLTYIYLGEVLKTYN	120
QY	137	NSVLNTLIMISIVMVTIFLVLYKRYCYKFINGWLINSLMLFLTYIYLGEVLKTYN	196	137 NSVLNTLIMISIVMVTIFLVLYKRYCYKFINGWLINSLMLFLTYIYLGEVLKTYN	196
QY	121	VMDYPTLLTWNFGAVGMYC1HWKGP1YVLOQAYLIMISALMAYF1KYLPEWSAWYL	180	121 VMDYPTLLTWNFGAVGMYC1HWKGP1YVLOQAYLIMISALMAYF1KYLPEWSAWYL	180
QY	197	VMDYPTLLTWNFGAVGMYC1HWKGP1YVLOQAYLIMISALMAYF1KYLPEWSAWYL	256	197 VMDYPTLLTWNFGAVGMYC1HWKGP1YVLOQAYLIMISALMAYF1KYLPEWSAWYL	256

Qy 241 LPPDPEME-----EDSYDFGEPSPYVEFPLTGYPG-----EBL---- 277
Db 309 VPKNPKNTQNTORAERETDGGSGNDRGSEWEARDSHUGPHRSTPESRAAVQEULSGSI 368
Qy 277 --EBBEERGVKGIGDGFIFYSVLYVGKAATAGDWNNTLACFVAAILGLCLTLIAVF 333
Db 369 LSEDEERGVKGIGDGFIFYSVLYVGKAATAGDWNNTIACFVAAILGLCLTLIAF 428
Qy 334 KKALPALPISITFGJLFYFESTDNLYVRPFENDTLASHOLYI 372
Db 429 KKALPALPISITFGJLFYFATDLYQPFMDQLAFHQFYI 467

RESULT 11
ID W23966 standard; Protein; 467 AA.
AC W23966;
DT 20-JUL-1998 (first entry)
DE Mouse Presenilin-1 homologue.
KW presenilin-1; PS1 gene; mouse; familial Alzheimer's disease; FAD; cerebral haemorrhage; schizophrenia; depression; epilepsy; mental retardation; diagnosis; therapy; transgenic animal.
KW protein; diagnosis; therapy; transgenic animal.
OS Mus musculus.
Key
FT Domain 82..100
FT Label= TM1
FT /note= "transmembrane domain 1"
FT Domain 101..132
FT Label= TM1-2
FT /note= "hydrophilic loop"
FT Domain 133..154
FT Label= TM2
FT /note= "transmembrane domain 2"
FT Domain 155..163
FT Label= TM2-3
FT /note= "hydrophilic loop"
FT Domain 164..183
FT Label= TM3
FT /note= "transmembrane domain 3"
FT Domain 184..194
FT Label= TM3-4
FT /note= "hydrophilic loop"
FT Domain 195..212
FT Label= TM4
FT /note= "transmembrane domain 4"
FT Domain 213..220
FT Label= TM4-5
FT /note= "hydrophilic loop"
FT Domain 221..238
FT Label= TM5
FT /note= "transmembrane domain 5"
FT Domain 239..243
FT Label= TM5-6
FT /note= "hydrophilic loop"
FT Domain 244..262
FT Label= TM6
FT /note= "transmembrane domain 6"
FT Domain 263..407
FT Label= TM6-7
FT /note= "hydrophilic loop"
FT Domain 408..428
FT Label= TM8
FT /note= "transmembrane domain 8"
FT Misc_difference 177
FT /note= "Phe177Ser mutation site (Claim 1)"
FT Misc_difference 439
FT /note= "Ile439Val mutation site (Claim 1)"
PN W09801549-A2.
PD 15-JAN-1998.
PF 04-JUL-1997; CA0475.
PT 02-JAN-1997; US-034590.
PR 05-JUL-1996; US-021673.
PR 12-JUL-1996; US-021700.

PR 08-NOV-1996; US-029895.
PA (HCSR) HSC RES & DEV LP.
PA (UTOR) UNTY TORONTO GOVERNING COUNCIL.
PI Fraser PE, Rommens JM, St George-Hyslop PH;
DR WPI: 98-28355/25.
DR N-PDB: V04668.
PT New isolated mutant presenilin-1 genes - useful for developing products for use in detection, diagnosis and therapy of Alzheimer's disease and for drug screening.
PT Disclosure; Page 199-200; 238pp; English.
PS This polypeptide comprises the murine presenilin-1 (PS1) homologue.
CC Its amino acid sequence was deduced from an isolated cDNA clone (see V04668). Mutations in the human PS1 and PS2 genes (see V04666-68) have been linked to the development in humans of forms of familial Alzheimer's disease (FAD). All amino acids that are mutated in analysed FAD pedigrees (see W3964) were conserved in the murine homologue. Use of the nucleic acids and proteins comprising or derived from presenilins can be made in screening and diagnosing FAD, identifying and developing therapeutics for treatment of FAD, and in producing cell lines and transgenic animals useful as models of FAD. Methods for identifying substances that bind to, or modulate the activity of a presenilin protein, and methods for identifying substances that affect the interaction of a presenilin-interacting protein with a presenilin protein are also disclosed.
SQ Sequence 467 AA;

Query Match 74.4%; Score 1431.5; DB 1; Length 467;
Best Local Similarity 71.28%; Pred. No. 4e-147;
Matches 284; Conservative 36; Mismatches 50; Indels 29; Gaps 4;

Qy 1 EELTURGYAKHVKVIMLEPVPTLCMVIVVATIKSRSVREYTEKNGQLIYPTTEDPSVGRILL 60
Db 71 EELTURGYAKHVKVIMLEPVPTLCMVIVVATIKSRSVSEYTRKDQLIYPTTEDPSVGRILL 130
Qy 61 NSVLNLTLIMISIVVATIFLVLYKTCRCKFTHGWLMSSMLFLFTTYIIGEVLTYN 120
Db 131 HSILNRAIMISIVVATILLLVLYKTCRCKYTHAWLISLLFFSFILGEVFTYN 190
Qy 121 VAMDYPTTLLTLYWNFGAVGMYC1HWKGPLVLYQQLYIMISALMAYTKLPPEWSAVIL 180
Db 191 VADYTTVALIILWNGVGMIAHWGPRLQQLYIMISALMAYTKLPPEWTANIL 250
Qy 181 GAIISVDLVALICPKGFLMLVETAKERNEPFLALYSSAMWTVYGMAKLDPSSQCMLQ 240
Db 251 AVISYVDLVALICPKGFLMLVETAKERNEPFLALYSSAMWLYVMMAEGDPEAQ--RR 308
Qy 241 LPYDPMEME-----EDSYDSFGEPSYVEFPPLTGYPG-----EEL---- 277
Db 309 VPKNPNTNQRAERETDGSGNDGGFSEEVQAQDSHLGHRSPEPSRAVQELSGSI 368
Qy 277 ---EEEERGKVKGDFDIFYSVLYGKAATGSDDNNTLACFVAILIGCLTLLIAVF 333
Db 369 LTSEDPERGYKLGLDFIVSVLGRASATASGDNTNTIACFVAILIGCLTLLIAF 428
Qy 334 KKALPALPISITFGJLFYFSTDNLYVRPFMDTLASHOLYI 372
Db 429 KKALPALPISITFGLYFYFATDLYQPFMDQLAFHQFYI 467

RESULT 12
ID W11839 standard; Protein; 467 AA.
AC W11839;
DT 07-MAY-1997 (first entry)
DE Human early onset Alzheimer's disease (EOAD) polypeptide.
KW Early onset Alzheimer's disease; EOAD; neurodegenerative disease; Homo sapiens.
OS Key
FT misc_difference 26..29
FT /note= "unidentified amino acid residues"

CC alternate splicing of the genomic DNA sequence'. W05762 represents the
 CC coding sequence for wild type human PS-2. The presenilins are a family of
 CC highly conserved integral membrane proteins with a common structural motif,
 CC common alternative splicing patterns, and common mutational hot spot
 CC regions. Mutations in PS genes are implicated in familial Alzheimer's
 CC disease (AD) and possibly other diseases such as cerebral haemorrhage,
 CC schizophrenia, depression etc., so detection of mutations in the DNA
 CC encoding these sequences can be used for diagnosis of these diseases.
 CC These proteins, or vectors that express them or containing antisense
 CC sequences, antibodies selective for mutant forms of these proteins (such
 CC as W05762) and modulators of PS gene expression are potentially useful
 CC for treatment of AD etc. Transgenic animals are useful as models for drug
 CC screening. The antibodies can also be used e.g. for affinity purification
 CC and in immunoassays.

Sequence 467 AA:

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Query Match 74.3%; Score 1429.5; DB 1; Length 467;
Best Local Similarity 72.3%; Pred. No. 6.6e-147;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;
Qy 1 EELTLKYGAKHVMILFVPTLCHNIVVATIKSVRETYEENKQIYTPFREDTPSGQRL 60
Db 71 EELTLKYGAKHVMILFVPTLCHNIVVATIKSVFTRDQIYTPFEDTEVGQRL 130
Qy 61 NSVLTNLIMISVYVMTIFLVLYKRYCYKFLWMLSLMFLFETYIYLGEVLKTYN 120
Db 131 HS1LNAAMISVIVMTIILVLYKRYCYKVIARLISLLEFFSTYLGVEVKYN 190
Qy 121 VAMDYPILLTLYWNFGAYGMVC1HWKGPLVLYQDYLIMISALVALFVKYLPEMSAWYL 180
Db 191 VADYITVALLINFGVYGMISMHWKGPLRQLQDYLIMISALVALFVKYLPEMTAWYL 250
Qy 181 GAISVYDLYAVLCPKGPLRMLVTAQERNEPFIALYSSAMWTVGMARLDPSSQGL- 240
Db 251 AVISVYDLYAVLCPKGPLRMLVTAQERNETLFLPALLISSTMWLVNMAEGDPEAQRRVS 310
Qy 240 -QLPYDPE-MEEDSYDSDFE--PSVPEFPPLTGYPG-----EEL----- 277
Db 311 KNSKYNAASTERSQDTVAENDOGFSEWEARDSHLGPHRSTPESRAVQELSSSILA 370
Qy 277 -EEEEEERYVKLGDFIYFSVLYGKAATGSGDNNTLACFVALLIGCLTLLLAVERKK 335
Db 371 GEDEEERYVKLGDFIYFSVLYGKASATAGSDNTLACFVALLIGCLTLLLAVERKK 430
Qy 336 ALPALPISITFGLIFYFSTDNLVRPEMDTLASHOLYYI 372
Db 431 ALPALPISITFGLIFYFSTDLYQPFMDQLAHQFYI 467

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RESULT 15
 W28507 standard; Protein: 407 AA.
 AC W28507;
 DT 07-DEC-1997 (first entry)
 DT Partial AD3 sequence.
 KW AD3; AD3IP; Alzheimer's disease; chromosome; misssegregation;
 KW presenilin; inhibitor; AD; trisomy 21.
 OS Homo sapiens.

Key Location/Qualifiers
 misc_difference 86
 /label= mutation
 /note= "M -> L"
 misc_difference 103
 /label= mutation
 /note= "H -> R"
 misc_difference 186
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 /note= "A -> E"
 misc_difference 226
 /label= mutation
 /note= "L -> V"
 misc_difference 350

FT	/label= mutation
FT	/note= "C -> Y"
FT	W09707213-A2.
FT	PD 27-FEB-1997;
FT	PF 15-AUG-1996; U13314.
FT	PR 16-AUG-1995; US-002448.
FT	(HARD) HARVARD COLLEGE.
FT	PA
FT	PI Li J, Potter H;
FT	DR WPI; 97-15297/15.
FT	DR N-PSDB; T87402.
FT	Identifying genes which cause chromosome misssegregation - useful for PT identifying causes of and treatments for diseases, e.g. Alzheimer's PT disease, cancer and ageing
PS Disclosure: Fig 1: 77pp; English.	
CC Identifying genes which cause improper chromosome segregation, CC screening for inhibitors of chromosome misssegregation and processes CC caused by genes encoding chromosome misssegregation promoters CC was exemplified using Alzheimer's disease. The sequences CC given in T87401 to T87426 can be used in the above methods.	
CC The five mutations indicated in the Features Table cosegregate CC with early-onset familial Alzheimer's disease. It is predicted CC that these mutations result in increased levels of cells with CC trisomy 21 in carriers of the mutation compared with non-carriers.	
SQ Sequence 407 AA:	

Query Match 74.3%; Score 1429.5; DB 1; Length 407;
Best Local Similarity 72.3%; Pred. No. 5.4e-147;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;
Qy 1 EELTLKYGAKHVMILFVPTLCHNIVVATIKSVRETYEENKQIYTPFREDTPSGQRL 60
Db 11 EELTLKYGAKHVMILFVPTLCHNIVVATIKSVFTRDQIYTPFEDTEVGQRL 70
Db 121 NSVLTNLIMISVYVMTIFLVLYKRYCYKFLWMLSLMFLFETYIYLGEVLKTYN 120
Qy 61 NSVLTNLIMISVYVMTIFLVLYKRYCYKFLWMLSLMFLFETYIYLGEVLKTYN 120
Db 131 VADYITVALLINFGVYGMISMHWKGPLRQLQDYLIMISALVALFVKYLPEMTAWYL 190
Db 131 VADYITVALLINFGVYGMISMHWKGPLRQLQDYLIMISALVALFVKYLPEMTAWYL 190
Qy 181 GAISVYDLYAVLCPKGPLRMLVTAQERNEPFIALYSSAMWTVGMARLDPSSQGL- 240
Db 191 GAISVYDLYAVLCPKGPLRMLVTAQERNEPFIALYSSAMWTVGMARLDPSSQGL- 240
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Db 191 AVISVYDLYAVLCPKGPLRMLVTAQERNETLFLPALLISSTMWLVNMAEGDPEAQRRVS 310
Qy 240 -QLPYDPE-MEEDSYDSDFE--PSVPEFPPLTGYPG-----EEL----- 277
Db 251 KNSKYNAASTERSQDTVAENDOGFSEWEARDSHLGPHRSTPESRAVQELSSSILA 370
Db 311 GEDEEERYVKLGDFIYFSVLYGKAATGSGDNNTLACFVALLIGCLTLLLAVERKK 335
Db 371 GEDEEERYVKLGDFIYFSVLYGKASATAGSDNTLACFVALLIGCLTLLLAVERKK 430
Qy 336 ALPALPISITFGLIFYFSTDNLVRPEMDTLASHOLYYI 372
Db 431 ALPALPISITFGLIFYFSTDLYQPFMDQLAHQFYI 467

Search completed: March 20, 2000, 05:31:21
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SUMMARIES								
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3	1923	100.0	448	2	US-08-592-541-137	Sequence 137, App		
4	1923	100.0	372	2	US-08-592-541-138	Sequence 138, App		
5	1907.5	99.2	447	2	US-08-875-972-29	Sequence 29, App		
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7	1429.5	74.3	467	2	US-08-967-101-134	Sequence 134, App		
8	1429.5	74.3	407	2	US-08-875-972-4	Sequence 4, App		
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17	1127.5	58.6	541	2	US-08-967-101-166	Sequence 166, App		
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25	88.5	4.6	3169	2	US-08-477-451-6	Sequence 3, App		
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ALIGNMENTS								
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RESULT 1	US-08-967-101-137	;	Sequence 137,	Application US/08967101	;			
		;	Patent No. 5840540		;			
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		;	APPLICANT: ROMMENS, JOANNA M		;			
		;	APPLICANT: FRASER, PAUL E		;			
		;	TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED TO ALZHEIMER'S DISEASE		;			
		;	NUMBER OF SEQUENCES: 183		;			
		;	CORRESPONDENCE ADDRESS:		;			
		;	STREET: High Street Tower - 125 High Street		;			
		;	CITY: Boston		;			
		;	STATE: Massachusetts		;			
		;	COUNTRY: U.S.A.		;			
		;	Z-PP: 02110		;			
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		;	ATTORNEY/AGENT INFORMATION:		;			
		;	NAME: Pitcher, Edmund R.		;			
		;	TELECOMMUNICATION INFORMATION:		;			
		;	TELEPHONE: (617) 248-7100		;			
		;	TELEFAX: (617) 248-7100		;			
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		;	Qy 61 NSVLNTLIMISIVVMMTIFLVLYKRCYKFKIGWLIMSSMLLFLFTYIYLGEVLTKYN 120		;			
		;	Db 137 NSVLNTLIMISIVVMMTIFLVLYKRCYKFKIGWLIMSSMLLFLFTYIYLGEVLTKYN 196		;			

Db 77 EELTKYGAHKVIMLFVPTLCMIVVATKSVERXTERNGQLIYTPTFEDTPSGQRUL 136
 Qy 61 NSVLNTLIMISVYVMTIFLVLYKRYCKFIFHGWLIMSSMLLFLFTYIYGLEVLTKN 120
 Db 137 NSVLNTLIMISVYVMTIFLVLYKRYCKFIFHGWLIMSSMLLFLFTYIYGLEVLTKN 196
 Qy 121 VAMDYPPLLTTWNFPLQYALIMISALMALVFKYLPEWSAWIL 180
 Db 197 VAMDYPPLLTTWNFPLQYALIMISALMALVFKYLPEWSAWIL 256
 Qy 181 GAIISYDYLAVLPKGPKPLRMLVETQAERNEPFPALIYSSAMWVTVGMAKLDPSSQALQ 240
 Db 257 GAIISYDYLAVLPKGPKPLRMLVETQAERNEPFPALIYSSAMWVTVGMAKLDPSSQALQ 316
 Qy 241 LPYDPMEEDSYDSFGEPSPYPEFPPLTGYPGEELEEERGVKLGLGDFIFYSVLGK 300
 Db 317 LPYDPMEEDSYDSFGEPSPYPEFPPLTGYPGEELEEERGVKLGLGDFIFYSVLGK 376
 Qy 301 AAATGSGDNTTLACFVALIGLCLTLLIAVFKKALPALPISITFGLIFYFSTDNLVRP 360
 Db 377 AAATGSGDNTTLACFVALIGLCLTLLIAVFKKALPALPISITFGLIFYFSTDNLVRP 436
 Qy 361 FMDTFLASHOLYI 372
 Db 437 FMDTFLASHOLYI 448

RESULT 4
 US-08-592-541-138
 ; Sequence 138, Application US/08592541
 ; GENERAL INFORMATION:
 ; APPLICANT: ST. GEORGE-HYSLOP, PETER H
 ; APPLICANT: ROMMENS, JOHANNA M
 ; APPLICANT: FRASER, PAUL E
 ; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 ; TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
 ; NUMBER OF SEQUENCES: 183
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: TESTA, HURWITZ & THIBEAULT
 ; STREET: High Street Tower - 125 High Street
 ; CITY: Boston
 ; STATE: Massachusetts
 ; COUNTRY: U.S.A.
 ; ZIP: 02110
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patentin Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/592,341
 ; FILING DATE:
 ; CLASSIFICATION: 800
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Pitcher, Edmund R.
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (617) 248-7000
 ; TELEFAX: (617) 248-7100
 ; INFORMATION FOR SEQ ID NO: 138:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 372 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ; US-08-592-541-138

Qy 1 EELTKYGAHKVIMLFVPTLCMIVVATKSVERXTERNGQLIYTPTFEDTPSGQRUL 60
 Db 1 EELTKYGAHKVIMLFVPTLCMIVVATKSVERXTERNGQLIYTPTFEDTPSGQRUL 60
 Qy 61 NSVLNTLIMISVYVMTIFLVLYKRYCKFIFHGWLIMSSMLLFLFTYIYGLEVLTKN 120
 Db 61 NSVLNTLIMISVYVMTIFLVLYKRYCKFIFHGWLIMSSMLLFLFTYIYGLEVLTKN 196
 Qy 121 VAMDYPPLLTTWNFPLQYALIMISALMALVFKYLPEWSAWIL 180
 Db 121 VAMDYPPLLTTWNFPLQYALIMISALMALVFKYLPEWSAWIL 256
 Qy 121 VAMDYPPLLTTWNFPLQYALIMISALMALVFKYLPEWSAWIL 180
 Db 121 VAMDYPPLLTTWNFPLQYALIMISALMALVFKYLPEWSAWIL 180
 Qy 181 GAIISYDYLAVLPKGPKPLRMLVETQAERNEPFPALIYSSAMWVTVGMAKLDPSSQALQ 240
 Db 181 GAIISYDYLAVLPKGPKPLRMLVETQAERNEPFPALIYSSAMWVTVGMAKLDPSSQALQ 240
 Qy 241 LPYDPMEEDSYDSFGEPSPYPEFPPLTGYPGEELEEERGVKLGLGDFIFYSVLGK 300
 Db 241 LPYDPMEEDSYDSFGEPSPYPEFPPLTGYPGEELEEERGVKLGLGDFIFYSVLGK 300
 Qy 301 AAATGSGDNTTLACFVALIGLCLTLLIAVFKKALPALPISITFGLIFYFSTDNLVRP 360
 Db 301 AAATGSGDNTTLACFVALIGLCLTLLIAVFKKALPALPISITFGLIFYFSTDNLVRP 360
 Qy 361 FMDTFLASHOLYI 372
 Db 361 FMDTFLASHOLYI 372

RESULT 5 972-29
 US-08-875-972-29
 ; Sequence 29, Application US/08875972
 ; Patent No. 5985564
 ; GENERAL INFORMATION:
 ; APPLICANT: Huntingdon Potter and Jinhue Li
 ; TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
 ; TITLE OF INVENTION: CHROMOSOME NON-DISJUNCTION
 ; NUMBER OF SEQUENCES: 29
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
 ; STREET: Two Militia Drive
 ; CITY: Lexington
 ; STATE: Massachusetts
 ; COUNTRY: USA
 ; ZIP: 02173-4799
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patentin Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/875,972
 ; FILING DATE: 08-AUG-97
 ; CLASSIFICATION: 435
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: US 60/002,448
 ; FILING DATE: 16-AUG-1995
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Granahan Esq., Patricia
 ; REGISTRATION NUMBER: 32,227
 ; REFERENCE/DOCKET NUMBER: HU95-03PA
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (781) 861-6240
 ; TELEFAX: (781) 861-9540
 ; INFORMATION FOR SEQ ID NO: 29:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 447 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS:
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: protein
 ; US-08-875-972-29

Query Match 100.0% Score 1923; DB 2; Length 372;
 Best Local Similarity 100.0% Pred. No. 1.7e-190;
 Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 S-08-967-101-134

Query Match 8
 Best Local Similarity 74.3%; Score 1429.5; DB 2; Length 467;
 Matches 28; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

Y 1 EELTIKYGAKHVKIMLFVPPYFILCMIVWATIKSVRYTEKNGOLIYDPEFTDPSVGORLL 60
 Y 71 EELTIKYGAKHVKIMLFVPPYFILCMIVWATIKSVRYTEKNGOLIYDPEFTDPSVGORLL 130

Y 61 NSVNTLIMISVIVMTIFLVLYKRCYRPIHGWLJMSMILLFLFIFTYIYLGEVTKTYN 120
 b 131 HSILNAIMISVIVMTILLLWLYKRCYRPIHGWLJMSMILLFLFIFTYIYLGEVTKTYN 190

Y 121 VADNPYPLLTWLNFGAVGMVCIHWKGPLVQAYLIMISALMATEVIKYLPESAWYV 180
 b 191 VADVTIVALLIWNGGGVGMISIHWKGPRLQAYLIMISALMATEVIKYLPESAWYV 250

Y 181 GAIISYDVLVATCPKGPLRMLVETAQERNPPIFPALEYSSAMWTVGMAKLDPSQGAL- 240
 b 251 AVISYDVLVATCPKGPLRMLVETAQERNTLPALIYSSSTMWLVNMAEGPPEAQRRVS 310

Y 240 -QLPYDPE-MEEDSYDSFGE--PSEPVFPEPLTCYGP-----BELL----- 277
 b 311 KNSKYNAAESTERESDTVAINDGGSSEEVAQRDQHGLPRSTPSPRAVQELSSILA 370

Y 277 -EEPERGYKLGDFIYFSVLYGRAATGSGDWNTTLACTVAILIGLCITLILLAYFKK 335
 b 371 GEDPERGYKLGDFIYFSVLYGRAATGSGDWNTTLACTVAILIGLCITLILLAYFKK 430

Y 336 ALPALPISITFGLIFYFSTDNLVRPMDTIAASHQWLI 372
 b 431 ALPALPISITFGLIFYFATDYLVQSFMDQLAHFQFYI 467

RESULT 8
 Sequence 4, Application US/08875972
 Patent No. 5985564

GENERAL INFORMATION:
 APPLICANT: Huntington Potter and Jinhue Li
 TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
 CHROMOSOME NON-DISJUNCTION
 NUMBER OF SEQUENCES: 29
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
 STREET: Two Militia Drive
 CITY: Lexington
 STATE: Massachusetts
 COUNTRY: USA
 ZIP: 02173-4799

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/875,972
 FILING DATE: 08-AUG-97
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 60/002,448
 FILING DATE: 16-AUG-1995
 NAME: Granahan Esq., Patricia
 REGISTRATION NUMBER: 32,227
 TELECOMMUNICATION INFORMATION:

PHONE: (781) 861-6240
 TELEFAX: (781) 861-9540
 INFORMATION FOR SEQ ID NO: 4:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 407 amino acids
 TYPE: amino acid
 TOPOLogy: linear
 MOLECULE TYPE: protein
 US-08-875-972-4

Query Match 74.3%; Score 1429.5; DB 2; Length 407;
 Best Local Similarity 72.3%; Pred. No. 1..e-139; Mismatches 33; Indels 25; Gaps 5;
 Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

QY 1 EELTIKYGAKHVKIMLFVPPYFILCMIVWATIKSVRYTEKNGOLIYDPEFTDPSVGORLL 60
 Db 11 EELTIKYGAKHVKIMLFVPPYFILCMIVWATIKSVRYTEKNGOLIYDPEFTDPSVGORLL 70

QY 61 NSVNLIMISVIVMTIFLVLYKRCYRPIHGWLJMSMILLFLFIFTYIYLGEVTKTYN 120
 Db 71 HSILNAIMISVIVMTILLLWLYKRCYRPIHGWLJMSMILLFLFIFTYIYLGEVTKTYN 130

QY 121 VADNPYPLLTWLNFGAVGMVCIHWKGPLVQAYLIMISALMATEVIKYLPESAWYV 180
 Db 131 VADVTIVALLIWNGGGVGMISIHWKGPRLQAYLIMISALMATEVIKYLPESAWYV 190

QY 181 GAIISYDVLVATCPKGPLRMLVETAQERNPPIFPALEYSSAMWTVGMAKLDPSQGAL- 240
 Db 191 AVISYDVLVATCPKGPLRMLVETAQERNPPIFPALEYSSAMWTVGMAKLDPSQGAL- 250

QY 240 -QLPYDPE-MEEDSYDSFGE--PSEPVFPEPLTCYGP-----BELL----- 277
 Db 251 KNSKYNAAESTERESDTVAINDGGSSEEVAQRDQHGLPRSTPSPRAVQELSSILA 310

QY 277 -EEPERGYKLGDFIYFSVLYGRAATGSGDWNTTLACTVAILIGLCITLILLAYFKK 335
 Db 311 GEDPERGYKLGDFIYFSVLYGRAATGSGDWNTTLACTVAILIGLCITLILLAYFKK 370

QY 336 ALPALPISITFGLIFYFSTDNLVRPMDTIAASHQWLI 372
 Db 371 ALPALPISITFGLIFYFATDYLVQSFMDQLAHFQFYI 407

RESULT 9
 US-08-592-541-134
 Sequence 134, Application US/08592541
 Patent No. 5986054

GENERAL INFORMATION:
 APPLICANT: ST. GEORGE-HYSLOP, PETER H
 APPLICANT: ROMMEN, JOHANNA M
 APPLICANT: FRASER, PAUL E
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 NUMBER OF SEQUENCES: 183
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: TESTA, HURWITZ & THIBEAULT
 STREET: High Street Tower - 125 High Street
 CITY: Boston
 STATE: Massachusetts
 COUNTRY: U.S.A.
 ZIP: 02110

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/875,972
 FILING DATE: 08-AUG-97
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 60/002,448
 FILING DATE: 16-AUG-1995
 NAME: Granahan Esq., Patricia
 REGISTRATION NUMBER: 32,227
 TELECOMMUNICATION INFORMATION:
 ATTORNEY/AGENT INFORMATION:
 NAME: Pittman, Edmund R
 NUMBER: HU95-03PA

OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/670,964
FILING DATE: 26-JUN-1996
CLASSIFICATION: 435
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 60/001,142
FILING DATE: 13-JUL-1995
APPLICATION NUMBER: 60/001,501
FILING DATE: 18-JUL-1995
ATTORNEY/AGENT INFORMATION:
NAME: Han, William T.
REGISTRATION NUMBER: 34,344
REFERENCE/DOCKET NUMBER: P50358
TELECOMMUNICATION INFORMATION:
TELEPHONE: 610-270-5219
TELEFAX: 610-270-5090
TELEX:
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 463 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-670-964-4

Query Match 74.3%; Score 1429.5; DB 3; Length 463;
Best Local Similarity 72.3%; Pred. No. 1.9e-139; Mismatches 33; Indels 25; Gaps 5;
Matches 287; Conservative 33; Mismatches 52; Indels 25; Gaps 5;

Qy 1 EELTKYGAKHVINLFPVPLCMIVVATIKSVRFYETENGQLIYTFEDTPSGQRLL 60
Db 67 EELTKYGAKHVINLFPVPLCMIVVATIKSVRFYETENGQLIYTFEDTPSGQRLL 126
Qy 61 NSVNLTLIMISVIVMTIFLVLYKRYCKFKIPIHGWLIMSSLMLFLFTYIYLGEVLTYN 120
Db 127 HSILNRAIMISVIVMTILLVVLYKRYCKVKVHAWLIISSLLUFFSFITYLGEVFTYN 186
Qy 121 YAMDYPPTLITVWNFGAVGMVCIHWKGPFLVLYQAYLIMISALMAYFIKYLPEWSAWVIL 180
Db 187 VADYTIVAILVNNFGVVGMISIHWKGPFLRQLQAYLIMISALMAYFIKYLPEWTAWLIL 246
Qy 181 GAIISYDLYAVLPKGLPKLMLVTAERNEPFPALIYSSAMWTVGMAKLDPSQGL - 240
Db 247 AVISYDLYAVLPKGLPKLMLVTAERNETFPALIYSSTMWLVNMAEGDPEAQRRVS 306
Qy 240 -QLPYDPE-MEEDSYDSFGE--PSYPEVEFPPLTGYPG-----EEL----- 277
Db 307 KNSKVNAAESTERESDVTENDGGFSEENEAQRDSHLGPHRSPSRAVEQELSSILA 366
Qy 277 -EEPEERGYVKLGDFIIFYSVLYGKAATGSDWNTTLACFVALIGCLTLLLAVFKK 335
Db 367 GEDPPERGVKLGLGDFFISVLYGRASATASGDWNTTIACFVALIGCLTLLLAIFKK 426
Qy 336 ALPALPISITFGFLIFYSTDNLVRPFMDTLASHQLYI 372
Db 427 ALPALPISITFGFLIFYFATDLYQPFMDQLAFHQFYI 463

RESULT 12
US-08-967-101-2
; Sequence 2, Application US/08967101
; Patent No. 5840540

GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H
; APPLICANT: ROMMENS, JOHANNA M
; APPLICANT: FRASER, PAUL E
; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
; TO ALZHEIMER'S DISEASE
; NUMBER OF SEQUENCES: 183

RESULT 13
US-08-592-541-2
; Sequence 2, Application US/08592541
; Patent No. 5986054
; GENERAL INFORMATION:
; APPLICANT: ST. GEORGE-HYSLOP, PETER H

RESULT 15
 US-08-967-101-4
 Sequence 4, Application US/08967101
 Patent No. 5840540

GENERAL INFORMATION:
 APPLICANT: FRASER, PAUL E
 ROMMEN, JOANNA M
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED TO ALZHEIMER'S DISEASE
 NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:
 ADDRESSEE: TESPA, HURWITZ & THIBEAULT
 STREET: High Street Tower - 125 High Street
 CITY: Boston
 STATE: Massachusetts
 COUNTRY: U.S.A.
 ZIP: 02110

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/967/101
 FILING DATE: 10-NOV-1997
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/592,541
 FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Pitcher, Edmund R.
 TELECOMMUNICATION INFORMATION:

TELEPHONE: (617) 248-7000

TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 4:

SEQUENCE CHARACTERISTICS:

LENGTH: 467 amino acids

STRANDEDNESS: single

TOPOLOGY: Linear

MOLECULE TYPE: protein

US-08-967-101-4

Db 367 GEDPERGVKLGLDFIFYSVLYGRASATASGDNNTTIACFVALLIGLCLTLIAFRK 426
 Qy 336 ALPALPISITFGCLFYFSTDNLVRPEMDTLASHQLYI 372
 Db 427 ALPALPISITFGCLFYFATDYLWQPMQLAHQFYI 463
 RESEQ 15
 US-08-967-101-4
 Sequence 4, Application US/08967101
 Patent No. 5840540

APPLICANT: ST. GEORGE-HYSLOP, PETER H
 ROMMEN, JOANNA M
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED TO ALZHEIMER'S DISEASE
 NUMBER OF SEQUENCES: 183

ADDRESSEE: TESPA, HURWITZ & THIBEAULT
 STREET: High Street Tower - 125 High Street
 CITY: Boston
 STATE: Massachusetts
 COUNTRY: U.S.A.
 ZIP: 02110

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/967/101
 FILING DATE: 10-NOV-1997
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/592,541
 FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Pitcher, Edmund R.
 TELECOMMUNICATION INFORMATION:

TELEPHONE: (617) 248-7000

TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 4:

SEQUENCE CHARACTERISTICS:

LENGTH: 467 amino acids

STRANDEDNESS: single

TOPOLOGY: Linear

MOLECULE TYPE: protein

US-08-967-101-4

Search completed: March 18, 2000, 19:55:32
 Job time: 3230 sec

Query Match 71.8%; Score 1381.5; DB 2; Length 467;
 Best Local Similarity 68.7%; Pred. No. 1.8e-134;
 Matches 274; Conservative 38; Mismatches 58; Indels 29; Gaps 4;

Qy 1 EELTIKYGAKHVNMLFPVPLCIVVVVATIKSPTREYTERNGQLIYTPFTEDTPSVQRLL 60
 Db 71 EELTIKYGAKHVNMLFPVPLCIVVVVATIKSPTREYTERNGQLIYTPFTEDTPSVQRLL 130

Qy 61 NSVLNTLIMISIVVMTIPLVYVLYKRYCYKFLFVWLQAYLIMISALMVFYLPEWSAWIL 120
 Db 131 HS1UNAAAMISIVVMTIPLVYVLYKRYCYKFLFVWLQAYLIMISALMVFYLPEWSAWIL 190

Qy 121 VAMDYPTLLLTVWNFGAYGMVCIHWKGPLVLYQAYLIMISALMVFYLPEWSAWIL 180
 Db 191 VVXYVTYVALLIWNWGVYGMIAHWKGPLRQAYLIMISALMVFYLPEWSAWIL 250

Qy 181 GAISSYDLYAVLCPFKPLMVLVTAERNEPIFALIYSSAMYWTVMGAKLDPSSQALQ 240
 Db 251 AVISYDLYAVLCPFKPLMVLVTAERNETLFALIYSSSTMVLVNAEGDPEAQ--RR 308

probable O-antigen
 cytochrome-c oxidase
 NADH dehydrogenase
 endothelin receptor
 oligopeptide transporter
 probable inner membrane
 probable membrane
 probable cell division
 glucose transporter
 bacteriochlorophyll
 GenCore version 4.5
 Copyright (c) 1993 - 1998 Compugen Ltd.
 Protein search, using sw model
 run on: March 18, 2000, 14:11:52 ; Search time 41.25 Seconds
 (without alignments)
 425,381 Million cell updates/sec

title: US-08-569-359B-138
rehearsal score 1923
sequence: 1 BELTLYGAKHIVMLFVBVT.....STDNLVRFMDTLAS HOLYI 372

scoring table: BLOSUM62
gapcost: 110000 20000 17160210 -20000

Number of hits that pass the threshold : 142080

2: pir2:*

4: Pir4: *

red. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution

IMMATURES

result No.	Description				
	Query Score	Match Length	DB ID	ID	
1	1923	100.0	448	2	I58098
2	1915	99.6	448	2	A56993
3	1802.5	93.7	442	2	I39174
4	1534	79.8	449	2	JC5391
5	1431.5	74.4	467	2	I78388
6	1431.5	74.4	433	2	JC5390
7	1429.5	74.3	467	2	S563683
8	1429.5	74.3	463	2	S63683
9	1413.5	73.5	463	2	JC5081
10	1413.5	73.5	467	2	JC5082
11	1011	52.6	374	2	S63684
12	956	49.7	461	2	S66253
13	524.5	27.3	358	2	T15184
14	513	26.7	453	2	T00724
15	274	14.2	465	2	A13459
16	110	5.7	2016	2	A31195
17	104	5.4	826	2	T02268
18	104	5.4	398	2	H75043
19	103	5.4	382	2	S41882
20	102	5.3	379	2	I48135
21	101.5	5.3	1681	2	A55138
22	100.5	5.2	324	2	S31646
23	99.5	5.2	1840	1	CHATM1
24	99.5	5.2	379	2	JC6178
25	99	5.1	2019	2	A313996
26	98	5.1	531	2	T11596
27	98	5.1	461	2	T11829
28	98	5.1	217	2	S01095
29	97	5.0	381	2	T11440
30	97	5.0	447	2	S52968
31	97	5.0	238	2	S02063
32	97	5.0	441	2	S11425
33	97	5.0	299	2	D65187
34	96.5	5.0	308	2	S22928
35	96	5.0	1020	2	S22929

ATTACHMENT S

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RESULT 1
I58038
E5-1 protein - human
C;Species: Homo sapiens (mam)
C;Accession: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 29-Sep-1999
C;Accession: I58098
C;Accession: I58098
R;Rogaeva, E.I.; Sherrington, R.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Liang, J.; Cohen, D.; Lannert, L.; Fraser, P.E.; Rommens, J.M.; St. George-Hyslop, P.H.
Nature 376, 775-778, 1995
A;Title: Familial Alzheimer's disease in kindreds with missense mutations in a

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A;Accession: I38098
A;Status: preliminary; translated from GB/EMBL/DDJB
A;Molecule type: mRNA
A;Residues: 1-448 <R>
A;Cross-references: GB:L44577; NID:950347; PTDN: AAC42012.1; PID: g950348

C; Genetics: A; Gene: E5-1
C; Superfamily: presenilin

Query	Match	Score	Length
QY	1 EELTIKYGAKHVMILFVPTVTLCMIVVVATIKSYSTREYTAKNGQLIYTPTEDTPSVQGRLL	100.0%	21
Best Local Similarity	Score 1923; DB 2;	100.0%;	Length 448;
Matches	Pred. No. 3.5e-138;	0;	
Matches	Mismatches	0;	Gaps
Db	77 EELTIKYGAKHVMILFVPTVTLCMIVVVATIKSYSTREYTAKNGQLIYTPTEDTPSVQGRLL	100.0%	136
QY	61 NSVNTLIMISVTVMTIFLVLYKRYCKFKFGWLINSSMLLFLFTTYIYGVLKYN	100.0%	120
Db	137 NSVNTLIMISVTVMTIFLVLYKRYCKFKFGWLINSSMLLFLFTTYIYGVLKYN	100.0%	196
QY	121 VANDYPTLLLTVWNFGAYGMVCYTHWKGPFLVLYQAYLIMISALMALVFKYLPENSANTYL	100.0%	180
Db	197 VANDYPTLLLTVWNFGAYGMVCYTHWKGPFLVLYQAYLIMISALMALVFKYLPENSANTYL	100.0%	256
QY	181 GAIISYDLYAVLCPKGPRMLVETAQNENEPIPALIYSSAMWVTGMAKLDDSSQGLQ	100.0%	240
Db	257 GAIISYDLYAVLCPKGPRMLVETAQNENEPIPALIYSSAMWVTGMAKLDDSSQGLQ	100.0%	316
QY	241 LPDYPEMEDDSFGEPSPYPEVEPPPLTGYPGEBLEEEERGGYKLGIGDFITYSVLYGK	100.0%	300
Db	317 LPDYPEMEDDSFGEPSPYPEVEPPPLTGYPGEBLEEEERGGYKLGIGDFITYSVLYGK	100.0%	376
QY	301 AAANGSDWNTTACFVALIGLCLTLLAAVKKALPAPISITGFLIFYFSTDNLWRP	100.0%	360

三

361 FMDTLASHOLYI 3/4
437 FMDTLASHOLYI 4/4

RESULT 2
A56993
Presentin 2 - human

Qy	61	NSYLTNLIMISVIVMTIFLVVLYKRCYKFIRGWLMSSLMLFLFTYIYLGEVLKTYN	120	Qy	181	GAIISYDLYAVLCPKGPLMLVETAERNPFPALIYSSAMWVTVGMAKLDPSOGAL-	240
Db	127	HSTLNAAAMISVIVMTIFLVVLYKRCYKFIRGWLMSSLMLFLFTYIYLGEVLKTYN	186	Db	251	AVISYDLYAVLCPKGPLMLVETAERNPFPALIYSSAMWVTVGMAKLDPSOGAL-	310
Qy	121	VADYPTPLTLLTVNFGAYGMVCIHWKGPVLVQAYLIMISALMALVFIFYLGEVKYN	180	Qy	240	-QLPYD-----PEMEEDSYDSFGEPSYPEVEP---PLTGYPGEEL---	277
Db	187	VADYITVALLINNGVYGMISHWKGPVLRLQQAYLIMISALMALVFIFYLGEVKYL	246	Db	311	KNTKYNAQCTEREAQASVPPNDGGFSEWEAQQRSDOLGHRSSTSVAEVOEISSIPA	370
Qy	181	GASVYDLYAVLCPKGPLMLVETAERNPFPALIYSSAMWVTVGMAKLDPSOGAL-	240	Qy	277	-EEEBERGKIGLGLDFEFYSLVYKRAAATASGDWNTTACFVAILIGLCTLILAYFK	335
Db	247	AVISYDLYAVLCPKGPLMLVETAERNETLPALEYSSSTMWLVNAAEGDBEAQRVVS	306	Db	371	SEDPERGTKLGLGDFEVTSVLYGRASATASGDWNTTACFVAILIGLCTLILAYFK	430
Qy	240	-QLPYD-----PEMEEDSYDSFGEPSYPEVEP---PLTGYPGEEL---	277	Qy	336	ALPALPISITFGLIFYFSTDNLVRPFMDTLASHQYI	372
Db	307	KNTKYNAQCTEREAQASVPPNDGGFSEWEARDSQLGPFRSTSVAEVOEISSIPA	366	Db	431	ALPALPISITFGLIFYFSTDNLVRPFMDQLAFHQFYI	467
Qy	277	-EEEBERGKIGLGLDFEFYSLVYKRAAATASGDWNTTACFVAILIGLCTLILAYFK	335	RESULT 11			
Db	367	SEDPERGTKLGLGDFEVTSVLYGRASATASGDWNTTACFVAILIGLCTLILAYFK	426	Qy	63684	presenilin 1, splice form 374 - human	
Qy	336	ALPALPISITFGLIFYFSTDNLVRPFMDTLASHQYI	372	C;Species: Homo sapiens (man)			
Db	427	ALPALPISITFGLIFYFSTDNLVRPFMDQLAFHQFYI	463	C;Date: 20-Jul-1996 #sequence_revision 13-Mar-1997 #text_change 29-Sep-1999			
RESULT 10				C;Accession: S6644			
JC5080		presenilin 1 protein isoform 467 - lesser mouse lemur		A;Status: preliminary; nucleic acid sequence not shown			
C;Species: Microcebus murinus (lesser mouse lemur)				A;Accession: S63684			
C;Date: 31-Jan-1997 #sequence_revision 31-Jan-1997				A;Title: Identification and characterization of presenilin I-467, I-463 and I-374.			
C;Accession: JC5080				A;Reference number: S63683; MUID:95193901			
R;Academia, Ai; Mestre Frances, N; Czech, C.; Pradier, L.; Petter, A.; Bons, N.; Bellis, Biochem. Biophys. Res. Commun. 248, 430-439, 1996				A;Cross-references: EMBL:U40380; PID:91244639; PIDN:AA05895.1; PID:91244640			
A;Title: Molecular cloning, sequencing, and brain expression of the presenilin 1 gene in				A;Gene: GDB:PSEN1; AD3: FAD; S182; PS1			
A;Reference number: JG3080; MUID:9707199				A;Cross-references: GDB:135682; OMIM:104311			
A;Accession: JG3080				A;Map position: 14q24.3-14q24.3			
A;Molecule type: mRNA				C;Superfamily: presenilin			
A;Residues: 1-467 <CAL>				Query Match 52.6%; Score 1011; DB 2; Length 374;			
C;Keywords: transmembrane protein				Best Local Similarity 81.8%; Pred. No. 2.4e-60;			
F;82-100/Domain: transmembrane #status predicted <TM1>				Mismatches 18; Indels 0; Gaps 0;			
F;133-154/Domain: transmembrane #status predicted <TM2>				Qy 1 EELTLKYGAKHVINMLFVPVTLCLMIVVVATIKSVRYTEKNGQLITPTPSVGQLL 60			
F;164-185/Domain: transmembrane #status predicted <TM3>				Db 67 FELTLKYGAKHVINMLFVPVTLCLMIVVVATIKSVRYTEKNGQLITPTPSVGQLL 126			
F;195-213/Domain: transmembrane #status predicted <TM4>				Qy 61 NSVINTLIMISVIVMTIFLVLYKRCYKFIRGWLMSSLMLFLFTYIYLGEVKYN 120			
F;221-238/Domain: transmembrane #status predicted <TM5>				Db 127 HSILNAAIMISVIVMTIFLVLYKRCYKFIRGWLMSSLMLFLFTYIYLGEVKYN 186			
F;244-261/Domain: transmembrane #status predicted <TM6>				Qy 121 VAMDYPTLILTVNFGAYGMVCIHWKGPLVQAYLIMISALMALVFIFYLPEWSAWVIL 180			
F;408-428/Domain: transmembrane #status predicted <TM7>				Db 187 VADYDITVALLINNGVYGMISHWKGPRLQQAYLIMISALMALVFIFYLPEWTAWIL 246			
Qy				Qy 181 GAIISYDLYAVLCPKGPLMLVETAERNPFPALIYSSAMWVTVGMAKLDPSOGAL 236			
Best Local Similarity 73.5%; Score 1413.5; DB 2; Length 467;				Db 247 AVISYDLYAVLCPKGPLMLVETAERNETLFPALIYSSAMWVTVGMAKLDPSOGAL 302			
Matches 282; Conservative 34; Mismatches 56; Indels 25; Gaps 4;				RESULT 12			
Qy	1	EELTLKYGAKHVINMLFVPVTLCLMIVVVATIKSVRYTEKNGQLITPTPSVGQLL	60	S6023	sel-12 protein - Caenorhabditis elegans		
Db	71	EELTLKYGAKHVINMLFVPVTLCLMIVVVATIKSVRYTEKNGQLITPTPSVGQLL	130	C;Species: Caenorhabditis elegans			
Qy	61	NSVINTLIMISVIVMTIFLVLYKRCYKFIRGWLMSSLMLFLFTYIYLGEVKYN	120	C;Date: 10-Apr-1996 #sequence_revision 19-Apr-1996			
Db	131	HSTLNAAAMISVIVMTIFLVLYKRCYKFIRGWLMSSLMLFLFTYIYLGEVKYN	190	C;Accession: S60253			
Qy	121	VADYPTPLTLLTVNFGAYGMVCIHWKGPLVQAYLIMISALMALVFIFYLPEWSAWVIL	180	R;Levitash, D.; Greenwald, I.			
Db	191	VADYITVALLINNGVYGMISHWKGPRLQQAYLIMISALMALVFIFYLPEWTAWIL	250	Nature 37, 351-354, 1995			
				A;Title: Facilitation of lin-12-mediated signalling by sel-12, a Caenorhabditis elega			
				A;Reference number: S60253; MUID:96032531			
				A;Accession: S60253			
				A;Status: preliminary; nucleic acid sequence not shown			

Db 369 KLGQDFIFTYSVNLVGRAAMY--DLMVTYACYLAIISGLGCTTLLSYNRAALPALPISI 425
 Qy 345 TFGLIFYFSTDNLVRFM 362
 Db 426 MLGVVFYFLTRLLMFPV 443

RESULT 15

A43459 sperm membrane protein spe-4 - Caenorhabditis elegans

N: Alternate names: probable integral membrane protein
C: Species: *Caenorhabditis elegans*

C: Date: 10-Jun-1993 #sequence 18-Nov-1994 #text_change 09-Sep-1997

C: Accession: A43459; S24632; S2433

R: L'Herault, S.W.; Arduengo, P.M.

J. Cell Biol. 119: 55-88. 1992

A: Title: Mutant of a putative sperm membrane protein in *Caenorhabditis elegans* prevent

A: Reference number: A43459; MUID:9240/040

A: Accession: A43459

A: Status: preliminary; not compared with conceptual translation

A: Molecule type: DNA; mRNA

A: Residues: 1-465 <LHE>

A: Cross-references: EMBL:Z14067; NID:96868; PID:96869; EMBL:Z14066; NID:96870; PID:96871

A: Experimental source: strain Bristol N2

A: Note: the nucleotide sequence was submitted to the EMBL Data Library, July 1992

C: Genetics:

A: Introns: 69/3; 154/3; 200/1; 224/3; 300/1; 385/1; 435/1

Query Match 14.2%; Score 274; DB 2; Length 465;
Best Local Similarity 21.5%; Pred. No. 1.5e-13;
Matches 93; Conservative 80; Mismatches 127; Indels 132; Gaps 14;

Qy 38 EKNGQLITYTPFTEDT--PSVGQRLILNSVLN---TLIMISIVYMTIFLVLYKYRCYKF 91

Db 42 EVNSELSKSTYFLDPSFQEQTGNLILDGFINGVTFILVUGCVSITMLAF--VLDFER--RI 97

Qy 92 IIGWULIMSSLMLF-----LFTYIYLGEVLTXYNADYPTILL----TWNFGA 137

Db 98 VKAWLTLSCLLIFGVSQTLHMFQSQFDQDNQX -----YMTIVLIVVPPVYGRG- 152

Qy 138 VGMVCIHMKGPVLWQAYLIMISALMVFIKLPEWSAWIAGAISYDLYAVLCPKG 197

Db 152 -GIAFFSNSSLLHQIYVTVNLSLISYIPLVPSKTIWFWVIVLFDLFVLPNGP 210

Qy 198 LRMVETQAERNPPIPALIYSSAMWVYGMAKLDPSSQG-----ALQPYDPE 246

Db 211 LKVQEKASDYSICVNLIMFSANEKRLTAGSNEETNEGEESTIRRVKQTEYYTKRE 270

Qy 247 MEEDSY-----DSF-----GDSYYPEPEPLTGPFEELEEEEE-- 282

Db 271 AQDDEFYQKIRRRAAINPDSVTEHSPLVEAPSPIELKEKNST----EELSDESDTS 326

Qy 282 -----R-----R-----R 282

Db 327 EISGGSSNLISSSDSSTTYSTSDISTAEECDQKEDDLVNSLNDNDKRPATADALNQGE 386

Qy 283 GYRIGLGLDFITYSVLYGKAATSGDWNTLACFVAILIGLCLTLILAVFKALPALPI 342

Db 387 VLRIGFGDEVFYSLIGQAASGCP--FAVISRALGILFGLVVTVESTEESTTPALPL 444

Qy 343 SITEGLIYFST 354

Db 445 PVCGTFCYFSS 456

Qy	121 VADYPTPLLTWVNGFAYGMVCIHWKGPLVLQDYLIMISALNALVFIKLPWSANVIL 180 : : : : : : : : : : : Db	KW	Transmembrane; Glycoprotein. POTENTIAL.
Db	200 IADYPTPLVNINFGANGMICHWKGGLQDYLIMISALNALVFIKLPWSAWIL 259 : : : : : : : : : : : Qy	FT	TRANSMEM 133 103 POTENTIAL.
Qy	181 GATSYVDLAVICPKPLMVLTAERNEPIPPALIYSSAMMTVGMAKLDPSQGAL- 240 : : : : : : : : : : : Db	FT	TRANSMEM 161 181 POTENTIAL.
Db	260 GATSYVDLAVICPKPLMVLTAERNEPIPPALIYSSAMMTVGNA- SATADGRMN 31.8 : : : : : : : : : : : Qy	FT	TRANSMEM 195 215 POTENTIAL.
Qy	240 -QLY---DPNEMEDSYDSFGPSYPEVPEPLTGYGEELBEEERVKLGDFIFY 294 : : : : : : : : : : : Db	FT	TRANSMEM 221 241 POTENTIAL.
Db	319 QQYHIDRNTPEGANSTVEDAETRIQ -----TQSNISSEDPDEEVRVKLGDFIFY 371 : : : : : : : : : : : Qy	FT	TRANSMEM 244 264 POTENTIAL.
Qy	295 SLYVKGAAATGSGDWNTTLACFYAILIGLCLTLLAYVKAFLALPISITFGLIFYST 354 : : : : : : : : : : : Db	FT	TRANSMEM 281 301 POTENTIAL.
Db	372 SLYVKGAAATGSGDWNTTLACFYAILIGLCLTLLAYVKAFLALPISITFGLIFYST 431 : : : : : : : : : : : Qy	FT	TRANSMEM 408 428 POTENTIAL.
Qy	355 DNLYRPEMDTLASHOLYI 372 : : : : : : : : : : : Db	FT	TRANSMEM 433 453 POTENTIAL.
Db	432 DNLYRPEMDTLASHOMYI 449 : : : : : : : : : : : Qy	FT	CARBOHYD 30 30 POTENTIAL.
Qy	432 DNLYRPEMDTLASHOMYI 449 : : : : : : : : : : : Db	FT	CARBOHYD 279 279 POTENTIAL.
Qy	295 SLYVKGAAATGSGDWNTTLACFYAILIGLCLTLLAYVKAFLALPISITFGLIFYST 354 : : : : : : : : : : : Db	FT	CARBOHYD 405 405 POTENTIAL.
Qy	SEQUENCE 467 AA; 52639 MW; 3AE1350D CRC32;	SEQUENCE	467 AA; 52639 MW; 3AE1350D CRC32;
Qy	Query Match 74.4%; Score 1431.5; DB 1; Length 467; Best Local Similarity 71.2%; pred. No. 4.2e-91; Matches 284; Conservative 36; Mismatches 50; Indels 29; Gaps 4;		
Qy	1 EELTKYGAHKVIMLFVPTLCMIVVVAVATIKSVREYTEKNGOLIYTPEFDTPSVQRL 60 : : : : : : : : : : : Db	Db	71 EELTKYGAHKVIMLFVPTLCMIVVVAVATIKSVFYTTRDGQLIYTPEFDTETVQRAL 130 : : : : : : : : : : :
Qy	61 NSVNTLIMISVIVTMITFLVLYKRYCFLFVGLIMSSMLFLFTYTYLGEVLKTYN 120 : : : : : : : : : : : Db	Db	131 HSILNAAMISVIVTMITFLVLYKRYCFLFVGLIMSSMLFLFTYTYLGEVLKTYN 190 : : : : : : : : : : :
Qy	121 VAMDYPTLITLWVNGAVMVCTHWKGPLQDYLQQAYLIMISALMALVFKYHLISSLLEFFSFYLVGEVKTYN 180 : : : : : : : : : : : Db	Db	191 VAVDVTVAUWNGFGVVMIAIRWKGPRLQQAYLIMISALMALVFKYLPENAWLIL 250 : : : : : : : : : : :
Qy	181 GAISSYDIAVLCRKGLPMVLFVAFERNEPFIALYSSAMVITYGMAKLDPSSQALQ 240 : : : : : : : : : : : Db	Db	251 AVISYDYLAVLCPKGPLMLVTAQERNETLFPALEYSTMVWLNMAEGDPEAQ--RR 308 : : : : : : : : : : :
Qy	241 LPYDPEME-----EDSYDSFGGPSPYBFPEPLTGPG-----EEL---- 277 : : : : : : : : : : : Db	Db	309 VPKPKYNTQRAEBTQDSGSGNDGGFEEWQRDSDGPHRSTPESRAVQELSGSI 368 : : : : : : : : : : :
Qy	277 ---EEEEEYCYKLGLGDFLFYSVLYGKAATGSDDWNTLACEVAILIGLCLTLLAYV 333 : : : : : : : : : : : Db	Db	369 LTSEPEEEYKVLGDFYSVLYGKSATASGDWNTLACEVAILIGLCLTLLAYF 428 : : : : : : : : : : :
Qy	334 KKALPALPISITFOLIYESTDNLYRPFDTLASHQLYI 372 : : : : : : : : : : : Db	Db	429 KKALPALPISITFOLIYESTDNLYRPFDLYQPEMDQLAHQFYI 467 : : : : : : : : : : :
RESULT	7		
PSNL_XENIA	ID_PSNL_XENIA	STANDARD:	PRT; 433 AA.
PSNL_XENIA	ID_PSNL_XENIA	SEQUENCE FROM N.A.	
PSNL_XENIA	ID_PSNL_XENIA	AC 012976;	
PSNL_XENIA	ID_PSNL_XENIA	DT 15-JUL-1999 (Rel. 38, Created)	
PSNL_XENIA	ID_PSNL_XENIA	DT 15-JUL-1999 (Rel. 38, Last sequence update)	
PSNL_XENIA	ID_PSNL_XENIA	DT 15-JUL-1999 (Rel. 38, Last annotation update)	
PSNL_XENIA	ID_PSNL_XENIA	DE PRESENTIN ALPHABETIC.	
PSNL_XENIA	ID_PSNL_XENIA	GN PS-ALPHA.	
PSNL_XENIA	ID_PSNL_XENIA	OS Xenopus laevis (African clawed frog).	
PSNL_XENIA	ID_PSNL_XENIA	OC Batrachia; Anura; Mesobatrachia; Pipidae; Xenopodinae;	
PSNL_XENIA	ID_PSNL_XENIA	OC Xenopus.	
PSNL_XENIA	ID_PSNL_XENIA	RN SEQUENCE FROM N.A.	
PSNL_XENIA	ID_PSNL_XENIA	RX PRT; 433 AA.	
PSNL_XENIA	ID_PSNL_XENIA	RT TISSUE=BRAIN;	
PSNL_XENIA	ID_PSNL_XENIA	RL MEDLINE; 97223465.	
PSNL_XENIA	ID_PSNL_XENIA	RA TSUJIMURA A., YASOJIMA K., HASHIMOTO-GONOH T.; "Cloning of Xenopus presenilin-alpha and -beta cDNAs and their differential expression in oogenesis and embryogenesis.", Biochem. Biophys. Res. Commun. 231:392-396 (1997).	
PSNL_XENIA	ID_PSNL_XENIA	CC -!- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE.	
PSNL_XENIA	ID_PSNL_XENIA	CC -!- MAY FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS.	
PSNL_XENIA	ID_PSNL_XENIA	CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.	
PSNL_XENIA	ID_PSNL_XENIA	CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.	
PSNL_XENIA	ID_PSNL_XENIA	CC This SWISS-Prot entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL Outstation - the European Bioinformatics Institute. There are no restrictions on its use by non profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).	
PSNL_XENIA	ID_PSNL_XENIA	CC This SWISS-Prot entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL Outstation - the European Bioinformatics Institute. There are no restrictions on its use by non profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).	
PSNL_XENIA	ID_PSNL_XENIA	CC EMBL; L21177; AAC2094.1; -.	
PSNL_XENIA	ID_PSNL_XENIA	CC DR MGI; MG1_1202717; PSEN1; -.	
PSNL_XENIA	ID_PSNL_XENIA	CC DR MGD; MG1_1207510; ABB72049.1; -.	
PSNL_XENIA	ID_PSNL_XENIA	CC DR PFAM; PF01080; Presenilin; 1.	

- Db** 191 VADYITVALLINFGVVGMIAHHKGPLRLQQAYLIMISALMALKYKLPETWTLIL 250
QY 181 GATSYDYLAVLCPKGPLRMLVETAOERNEPIFPALIYSSAMVYGMALKDPSQGALQ 240
Db 251 AVISYDYLAVLCPKGPLRMLVETAOERNEPIFPALIYSSTMVLYNMAEGDPEAQ--RR 308
Db 241 LPYDP --- -EMEEDSYDSFGEPSYPVPEPPLTGYPG- -----EBL--- 277
Qy 309 VPKNPKYSTQSTEREFDTGSDGGFQHSHGPSTPESRAAQLSLSGS 368
Db 277 --- -EDEBERGVKLGLGDFIFYSVLYGKAATGSDWNNTLACFVAILIGLCLTLLAY 332
Qy 369 ILTSEDEPEEVGVKLGLGDFIFYSVLYGKAATGSDWNNTLACFVAILIGLCLTLLAY 428
Qy 333 FKKALPALPISITFLIFYFSTDNLYRPENDTLASHQLYI 372
Db 429 FKKALPALPISITFLIFYFSTDNLYQPFDOLAFHQFYI 468
- RESULT 9**
PSNL_HUMAN
 ID PSNL_HUMAN STANDARD; PRT; 467 AA.
 AC P49768; Q15762; Q15719; Q15720;
 DT 01-OCT-1995 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 15-JUL-1999 (Rel. 38, Last annotation update)
 DE PRESENILIN 1 (PS-1) (S182 PROTEIN).
 GN PSEN1 OR PSNL OR AD3 OR PS1 .
 OS Homo (Human).
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OC [1]
 RN SEQUENCE FROM N.A.; AND VARIANTS AD (FORMS I-467 AND I-463).
 RC TISSUE-BRAIN;
 RX MEDLINE; 95319502.
 RA SHERRETTING R., ROGAEV E.-I., LIANG Y., ROGAeva E.A., LEVESQUE G.,
 RA IKEDA M., CHI H., LIN C., LI G., HOLMAN K., TSUDA T., MAR L.,
 RA FONCINI J.-F., BRUNI A.C., MONTESI M.P., SORBI S., RAINERO L.,
 RA PINESI L., NEE L., CHUMAKOV I., POLLINI D., BROOKS A.,
 RA SANSEAU P., POLINSKY R.J., WASCO W., DA SILVA H.A.R., HAINES J.L.,
 RA PERICAS-VANCE M.A., VANCE R.E., TANZI R.E., ROSES A.D., FRASER P.E.,
 RA ROMMENS J.M., ST GEORGE-HYSLOP P.H.;
 RT "Cloning of a gene bearing missense mutations in early-onset familial Alzheimer's disease.";
 RT Nature 375:754-760(1995).
 RN [2] SEQUENCE FROM N.A. (FORMS I-463 AND I-374).
 RC TISSUE-BLOOD AND BRAIN;
 RX MEDLINE; 96193901.
 RA SHIRASAWA T., YAHAGI Y.-I., TAKAGI H., KONDO T., OKUCHI M., USAMI M.,
 RA RT "Identification and characterization of presenilin 1-467, I-463 and
 RT I-374.";
 RT FEBS Lett. 381:7-11(1996).
 RN [3] SEQUENCE OF 1-113 FROM N.A.
 RP TSUJIMURA A., HASHIMOTO-GOTO T.;
 RL Submitted (MAR-1996) to the EMBL/GenBank/DDBJ databases.
 RN [4] REVIEW ON VARIANTS.
 RX MEDLINE; 98180715.
 RA CRUTS M., VAN BROECKHOVEN C.;
 RT "Presenilin mutations in Alzheimer's disease.";
 RL Hum. Mutat. 11:183-190(1998).
 RN [5] VARIANTS AD THR-143 AND ALA-384.
 RX MEDLINE; 96177673.
 RA CRUTS M., BACKHOVEN H., WANG S.-Y., VAN GASSEN G., THEUNS J.,
 RA DE JONGHE C., WEHNERT A., DE VOECHT J., DE WINTER G., CRAS P.,
 RA BRUYLAND M., DATSON N., WEISSENBACH J., DEN DUNNEN J.T., MARTIN J.-J.,
 RA HENDRIKS L., VAN BROECKHOVEN C.;
 RT "Molecular genetic analysis of familial early-onset Alzheimer's
 disease linked to chromosome 14q24.3.";
- RL** Hum. Mol. Genet. 4:2363-2372(1995).
RN [6] VARIANTS AD I-82; H-115; T-139; R-163; T-231; L-264; V-392 AND Y-410.
RP MEDLINE; 96177674.
RA CAMPION D., FLAMAN J.-M., BRICE A., HANNEQUIN D., DUBOIS B.,
RA MARTIN C., MOREAU V., CHARBONNIER F., DIDIERJEAN O., TARDIEU S.,
RA PENET C., PUEL M., PASQUIER F., LE DOZE F., BELLIS G., CALENDRA A.,
RA HEILIG R., MARTINEZ M., MALLET J., BELLIS M., CLERGET-DARPOUX F.,
RA AGID Y., FREBOURG T.;
RT "Mutations of the presenilin 1 gene in families with early-onset Alzheimer's disease.";
RL Hum. Mol. Genet. 4:2313-2377(1995).
RN [7] VARIANTS AD VAL-260; VAL-285 AND VAL-392.
RP MEDLINE; 95379971.
RA ROGAEV E.I., SHERRETTING R., ROGAeva E.A., LEVESQUE G., IKEDA M.,
RA LIANG Y., CHI H., LIN C., HOLMAN K., MAR L., SORBI S.,
RA NACMIAS B., PIACENTINI S., AMADUCCI L., CHUMAKOV I., COHEN D.,
RA LANNEFELT L., FRASER P.E., ROMMENS J.M., ST GEORGE-HYSLOP P.H.;
RT "Familial Alzheimer's disease in kindreds with missense mutations in a gene on chromosome 1 related to the Alzheimer's disease type 3 gene.";
RT Nature 376:775-778(1995).
RN [8] VARIANTS AD V-139; V-146; Y-163; T-267; A-280 AND G-280.
RP MEDLINE; 96024664.
RA CLARK R.F., HUTTON M., FULLNER R.A., FROELICH S., KARRAN E.,
RA CLARK R.F., HUTTON M., FULLNER R.A., PRIHAR G., HE C., KORENBROT K.,
RA MARTINEZ A., CROOK R., LONDON C., PRIHAR G., HE C., KORENBROT K.,
RA MARTINEZ A., WRAGG M., BUSFIELD F., BEHRENS M., MYERS A., NORTON J.,
RA MORRIS J., MEITA N., PEARSON C., LINCOLN S., BAKER M., DUFF K.,
RA ZEHR C., PEREZ-TUR J., HOUDLEN H., RUIZ A., OSSA J., LOPERA F.,
RA ARCOS M., MADRIGAL L., COLLINGE J., HUMPHREYS C., ASWORTH T.,
RA SARNER S., FOX N., HARVEY R., KENNEDY A., ROQUES P., CLINE R.T.,
RA PHILLIPS C.A., VENTER J.C., FOREL L., AXELMAN K., LILIUS L.,
RA JOHNSTON J., COBBURN R., VIITANEN M., WINBLAD B., KOSIK K., HALTIA M.,
RA POYHONEN M., DICKSON D., MANN D., NEARY D., SNOWDEN J., LANTOS P.,
RA LANNEFELT L., ROSSOR M., ROBERTS G.W., ADAMS M.D., HARDY J., GOATE A.;
RT "The structure of the presenilin 1 (S182) gene and identification of six novel mutations in early onset AD families. Alzheimer's Disease Collaborative Group";
RT Neurosci. Lett. 208:195-198(1996).
RL Nat. Genet. 11:219-222(1995).
RN [9] VARIANTS AD PHE-96; ARG-163 AND THR-213.
RP MEDLINE; 96310408.
RA RAMINO K., SATO S., SAKAKI Y., YOSHIIWA A., NISHIWAKI Y., TAKEDA H.,
RA TANABE H., NISHIMURA T., LI K., ST GEORGE-HYSLOP P.H., MIKI T.,
RA OGIHARA T.;
RT "Three different mutations of presenilin 1 gene in early-onset Alzheimer's disease families.";
RL Neurosci. Lett. 208:195-198(1996).
RN [10] VARIANTS AD ALA-280.
RP MEDLINE; 97442268.
RA LONDON C.L., MARTINEZ A., BEHRENS I.M., KOSIK K.S., MADRIGAL L.,
RA NORTON J., NEUMAN R., MYERS A., BUSFIELD F., WRAGG M., ARCOS M.,
RA ARANGO VIANA J.C., OSSA J., RUIZ A., GOATE A.M., LOPERA F.;
RT "E280A PS-1 mutation causes Alzheimer's disease but age of onset is not modified by APOE alleles.";
RL Hum. Mutat. 10:186-195(1997).
RN [11] VARIANTS AD THR-233 AND THR-278.
RP MEDLINE; 97316242.
RA KWOK J.B.J., TADDEI R., HALLJUPP M., FISHER C., BROOKS W.S., BROE G.A.,
RA HARDY J., FULHAM M.J., NICHOLSON G.A., STELLI R.,
RA ST GEORGE-HYSLOP P.H., FRASER P.E., KAKULAS B., CLARNETTE R.,
RA RELKIN N., GANDY S.E., SCHOFIELD P.R., MARTINS R.N.;
RT "Two novel (M233T and R278T) presenilin-1 mutations in early-onset Alzheimer's disease pedigrees and preliminary evidence for association of presenilin-1 mutations with a novel phenotype.";
RL NeuroReport 8:1537-1542(1997).
RN [12] VARIANT GLY-318.
RP MEDLINE; 99115106.

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CC	EMBL; Z71333; CAA5930.1; -;	RT homologue.";
CC	DR; PF1080; Presenilin; 1.	RT NeuroReport 8:1025-1029(1997).
CC	KW Transmembrane; Alternative splicing; Glycoprotein.	RN [2]
CC	FT TRANSMEM 83 103	RP SEQUENCE FROM N.A. (SHORT ISOFORM).
CC	FT TRANSMEM 133 153	RC STRAIN=CANTON-S;
CC	FT TRANSMEM 161 181	RA HONG C.S., KOO E.H.;
CC	FT TRANSMEM 191 211	RT "Isolation and characterization of Drosophila presenilin homolog.";
CC	FT TRANSMEM 221 241	RA YE.Y., FORTINI M.E.;
CC	FT TRANSMEM 244 264	RT "Characterization of Drosophila presenilin and its colocalization with Notch during development.";
CC	FT TRANSMEM 281 301	RT Submitted (AUG-1998) to the EMBL/GenBank/DDBJ databases.
CC	FT TRANSMEM 408 428	RL Submitter (AUG-1998) to the EMBL/GenBank/DDBJ databases.
CC	FT TRANSMEM 433 453	CC -1 SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
CC	FT CARBOHYD 279 279	CC -1 SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC	FT CARBOHYD 405 405	CC -1 SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.
CC	FT VARSPPLIC 26 29	CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (see http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).
CC	SQ SEQUENCE 467 AA; 52384 MW; A841AOB7 CRC32;	CC DR EMBL; U77334; AAB61139.1; -;
CC	Query Match Score 1413.5; DB 1; Length 467; Best Local Similarity 71.0%; Pred. No. 7.2e-90; Matches 282; Conservative 34; Mismatches 56; Indels 25; Gaps 4;	CC DR EMBL; U78084; AAB53269.1; -;
Db	71 EELTKYGAHKVIMLFVPTLCLMIVVATIKSVRFYTYEKNGOLIYTPFTEDTPSGVORL 60	CC DR EMBL; AF04184; AAC33129.1; -;
Db	71 VADPYPTLLTWNNFGAVGMVCHEWKPLVLYQDYLIMTSALMALKFVLPENSAWIL 130	CC DR EMBL; AF04184; AAC33128.1; -;
Qy	61 NSVNTLIMISVIVMTIFLVVLYKCYRKFKFHGWLIMSSLMLFLFLTYVYLGEVLKRYN 120	CC DR FLYBASE; FBgn0019947; PS.
Db	131 HSYVNAATAMISVIVMVTILLVLYKCYRKVHAWLIISSLLFFSTYGEVFKTYN 190	CC DR PFAM; PF01080; Presenilin; 1.
Qy	121 VADPYPTLLTWNNFGAVGMVCHEWKPLVLYQDYLIMTSALMALKFVLPENSAWIL 180	CC KW Transmembrane; Glycoprotein; Alternative splicing.
Db	191 VAWXITYALLWNFGVGMISHEWKGPURLOQYLMISALMALKFVLPENTAWIL 250	CC FT TRANSMEM 107 127
Qy	181 GAISVSYDLYAVLCPKGPKPLRMLVYTAQERNEPPIFLAYSSAMWTVGMAKLDPSQGAL- 240	CC FT TRANSMEM 155 175
Db	251 AVISYDLYAVLCPKGPKPLRMLVYTAQERNETFLPALIYSTTMVLNNAEGDPAQRVVS 310	CC FT TRANSMEM 183 203
Qy	240 -QLPYD-----PEMEDSYDSFGEPSPYVEFEP---PLTGYPGEL----- 277	CC FT TRANSMEM 217 237
Db	311 KNTKYNAGQTERAQASVENDGGFSEWEAORDSOLGPHRSTSVAQOESSIPA 370	CC FT TRANSMEM 243 263
Qy	277 -EEBBERRYKLGLDFIFIVSVLKGKAATGSDWNTLACFATLIGCUTLILAVTRK 335	CC FT TRANSMEM 266 286
Db	371 SEDPPERGYKLGLDFEVYSLKGASAATGSDWNTLACFATLIGCUTLILAIFFK 430	CC FT TRANSMEM 304 324
Qy	336 ALPALPISITFGLLIFYFSTDNLVRFPEMDTLASHOLYI 372	CC FT TRANSMEM 482 502
Db	431 ALPALPISITFGLLDFEVYSLKGASAATGSDWNTLACFATLIGCUTLILAIFFK 467	CC FT TRANSMEM 507 527
Qy	1 EELTKYGAHKVIMLFVPTLCLMIVVATIKSVRFYTYEKNGOLIYTPFTEDTPSGVORL 60	CC FT TRANSMEM 527 547
Db	93 EEOQLKYGAHQVKLKFVPSLCLMIVVATINSYNTDVLLTPHEOSPEPSVKW 152	CC FT CARBOHYD 129 129
Qy	61 NSVNLIMISVIVMTIFLVVLYKCYRKFKFHGWLIMSSLMLFLFLTYVYLGEVLKRYN 120	CC FT CARBOHYD 339 339
Db	153 SALANSLIMMSVVVMTFLILVLLKRCYRTHGWLILSSFMLIFTYLYLEELRAYN 212	CC FT CARBOHYD 410 410
Qy	121 VAMDYPTLLTWNNFGAVGMVCHEWKPLVLYQDYLIMTSALMALKFVLPENSAWIL 180	CC FT CONFLICT 384 397
Db	213 IPMDYPTLLTWNNFGAVGMVCHEWKPLVLYQDYLIMTSALMALKFVLPENSAWIL 180	CC SQ MISSING (IN SHORT ISOFORM).
Qy	181 GAISYDLYAVLCPKGPKPLRMLVYTAQERNEPPIFLAYSSAMWTVGMAKLDPSQGAL- 240	CC FT CONFLICT 80 81
Db	273 AAISWDLIAVLSPRGPLRMLVYTAQERNEPPIFLAYSSAMWTVGMAKLDPSQGAL- 240	CC SQ GG -> RS (IN REF. 2).
Qy	231 LDPS-----QGALQLP-----YDPMEEDSYDSFGEPSPYEV-FEP 266	CC SQ 59304 MW; 79564FE0 CRC32;
Db	333 SSPSSNSITTRQNSLASPEAAAASGORTGNSHPRQNRDDGSVLTATEGMPVLFKS 392	CC SQ 541 AA; 58 6%; Score 1127.5; DB 1; Length 541;
Qy	1 EELTKYGAHKVIMLFVPTLCLMIVVATIKSVRFYTYEKNGOLIYTPFTEDTPSGVORL 60	CC Best Local Similarity 50.8%; Pred. No. 2.9e-70; Mismatches 61; Indels 74; Gaps 7;
Db	93 EEOQLKYGAHQVKLKFVPSLCLMIVVATINSYNTDVLLTPHEOSPEPSVKW 152	CC Matches 231; Conservative 231; Query Match 58 6%; Score 1127.5; DB 1; Length 541;

RESULT 11
 PSN_DROME STANDARD PRT; 541 AA.
 ID PSN_DROME STANDARD PRT; 541 AA.
 AC 002194; 002395; 076802;
 DT 15-JUL-1999 (Rel. 38, Created)
 DT 15-JUL-1999 (Rel. 38, Last sequence update)
 DT 15-JUL-1999 (Rel. 38, Last annotation update)
 DE PRESENILIN HOMOLOG (DPS) (DPS).
 PS Drosophila melanogaster (Fruit fly).
 OS Drosophila; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydriidae; Drosophilidae; Drosophila;
 RN [1] Sequence from N.A. (LONG ISOFORM).
 RX MEDLINE; 9728-868.
 RA BOUJANNE G.L., LIVNE-BAR I., HUMPHREYS J.M., LIANG Y., LIN C.,
 RA ROGAEV E., ST GEORGE-HYSLOP P.;
 RT "Cloning and characterization of the Drosophila presenilin

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OM protein - protein search, using sw mode.

Run on: March 18, 2000, 22:07:43 ; search time 49.26 Seconds
(without alignments)
523.595 Million cell updates/sec

Title: US-08-509-359B-138

Perfect score: 1923 Sequence: 1 EELTLYGAKHVINLFPVPT.....STDNLVRPFMDTLASHQLYI 372

Scoring table: BLOSUM62

Searched: 225878 seqs, 69334122 residues

Database : SPTREMBL_12; *

Word size : 0

Number of hits that pass the threshold : 225878

1: sp_archaea: * 2: sp_bacteria: * 3: sp_fungi: * 4: sp_human: * 5: sp_invertebrate: * 6: sp_mammal: * 7: sp_mhc: * 8: sp_organelle: * 9: sp_phage: * 10: sp_plant: * 11: sp_rabbit: * 12: sp_virus: * 13: sp_vertebrate: * 14: spUnclassified: *

Number of hits that pass the threshold : 225878

225878

ALIGNMENTS

Number	Score	Match	Length	DB	ID
26	103	5.4	266	8	Q34429 echimys did
27	103	5.4	266	8	Q37099 echimys did
28	103	5.4	380	8	Q9XNM7 microtus lo
29	102.5	5.3	382	8	Q34973 microtus d
30	102.5	5.3	382	8	Q35075 mastomys rub
31	102.5	5.3	382	8	Q35076 microtus r
32	102	5.3	444	2	Q85080 arthrobacte
33	102	5.3	266	8	Q34425 echimys did
34	102	5.3	266	8	Q34427 echimys did
35	102	5.3	379	8	Q34424 echimys did
36	102	5.3	379	8	Q33940 akodon uric
37	102	5.3	381	8	Q9XP85 smanthopsis
38	101.5	5.3	1681	11	Q62467 mus musculus
39	101	5.3	381	8	Q9XP88 smanthopsis
40	101	5.3	380	8	Q9XNL9 ondatra zib
41	100.5	5.2	382	8	Q35518 phialander m
42	100.5	5.2	324	11	Q61735 mus musculus
43	100	5.2	380	8	Q9XKB9 phaethon le
44	99.5	5.2	379	5	Q93127 balanus amp
45	99.5	5.2	382	8	Q355558 phialander o

ALIGNMENTS

RESULT	1
RESULT	1
Q9X96	PRELIMINARY;
ID	PRT;
Q9X96;	PRT;
AC	449 AA.
DT	01-NOV-1999 (TREMBLEUR. 12, Created)
DT	01-NOV-1999 (TREMBLEUR. 12, Last sequence update)
DT	01-NOV-1999 (TREMBLEUR. 12, Last annotation update)
DE	PRESENILIN 2.
OS	Bos taurus (Bovine).
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;
OC	Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovinae; Bos.
RN	[1]
RP	SEQUENCE FROM N.A.
RC	TISSUE=BRAIN;
RA	SAHARA N., SHIRASAWA T., MORI H.;
RT	"Molecular cloning of bovine presenilin 2 gene.";
RL	Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.
DR	EMBL: AF03893; Aad39024.1;
SQ	SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result	No.	Score	Query Match	Length	DB	ID	Description	Query Match	Length	DB	No.	Score	Best Local Similarity	Mismatches	Indels	Gaps
1	1896	98.6	449	6	Q9X96	Q9X96 bos taurus	1	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	120	Q9X96	98.6;	Score 1896;	DB 6;	Length 449;		
2	1411	73.4	478	6	Q9XT97	Q9XT97 bos taurinus	2	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	60	Q9X96;	98.4%	Pred. No. 9.9e-131;				
3	1388	72.7	456	13	Q9W677	Q9W677 brachydano	3	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	98.2%	Conservative 2;	Mismatches 4;			
4	1355	70.5	384	13	Q73859	Q73859 cyprinus ca	4	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	197	Q9X96;	98.1%	"Molecular cloning of bovine presenilin 2 gene."				
5	576	30.0	272	4	Q96340	Q96340 drosophila	5	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	98.0%	Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.				
6	402	20.9	184	4	Q95465	Q95465 homo sapiens	6	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.9%	DR AF03893; Aad39024.1;				
7	113	5.5	405	5	Q19737	Q19737 caenorhabditis	7	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.8%	SQ SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;				
8	112	5.8	4578	13	Q42281	Q42281 fugu rubrip	8	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.7%	RC TISSUE=BRAIN;				
9	110	5.7	320	8	Q34086	Q34086 coelcyzus er	9	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.6%	RA SAHARA N., SHIRASAWA T., MORI H.;				
10	110	5.7	381	8	Q35425	Q35425 phascolosior	10	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.5%	RT "Molecular cloning of bovine presenilin 2 gene."				
11	108	5.6	380	8	Q9ZC9	Q9ZC9 upupa epops	11	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.4%	RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.				
12	107	5.6	748	2	Q92577	Q92577 streptomyces	12	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.3%	DR AF03893; Aad39024.1;				
13	107	5.6	381	8	Q33723	Q33723 antechinus	13	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.2%	SQ SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;				
14	106	5.5	382	8	Q34340	Q34340 didelphis marsupialis	14	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.1%	RC TISSUE=BRAIN;				
15	106	5.5	318	11	Q3594	Q3594 rattus norvegicus	15	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	97.0%	RA SAHARA N., SHIRASAWA T., MORI H.;				
16	106	5.5	303	11	P97329	P97329 rattus norvegicus	16	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.9%	RT "Molecular cloning of bovine presenilin 2 gene."				
17	106	5.5	652	5	Q93346	Q93346 caenorhabditis	17	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.8%	RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.				
18	106	5.5	382	8	Q34279	Q34279 didelphis marsupialis	18	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.7%	DR AF03893; Aad39024.1;				
19	106	5.5	382	8	Q34677	Q34677 glironia ve	19	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.6%	SQ SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;				
20	105	5.5	318	11	Q9XN3	Q9XN3 arthrobacter	20	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.5%	RC TISSUE=BRAIN;				
21	105	5.5	379	8	Q34428	Q34428 echimys did	21	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.4%	RA SAHARA N., SHIRASAWA T., MORI H.;				
22	104	5.4	382	8	Q35561	Q35561 phialander o	22	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.3%	RT "Molecular cloning of bovine presenilin 2 gene."				
23	104	5.4	379	8	Q36096	Q36096 trinomys pa	23	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.2%	RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.				
24	104	5.4	379	8	Q34430	Q34430 echimys did	24	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.1%	DR AF03893; Aad39024.1;				
25	104	5.4	826	10	O80739	O80739 arabiopsis thaliana	25	EELTLYGAKHVINLFPVTLCKMIVVATIKSVRFYENKOLITYTPFTEDPSYQQLL	137	Q9X96;	96.0%	SQ SEQUENCE 449 AA; 50301 MW; A3DA878F CRC32;				

1	IMLFIPVTLCKMVVVATIKSVSFTYQDQQLYTPFREDTETVGQRANSMLNPAAMIS	60
72	VIVVMTIFLVLVLYKVRCKFIHGWMIMSSMLIPLIFTYIYLGEVILKTNYAMDYPTLLT	131
61	VIVVMTFLVLVLYKRCVKGWLFFSNMLLFFSFYLGEGEYKTNYAMDFETLAVI	120
132	WNNFGAVGMVICHWKGPVLQYALIMISALMAYFIKLPEWSAVLGIASTYDLVAV	191
121	IWNNGVGMICHWKGPRLRQYALIMISALMAYFIKLPEWTAWLILAASVYDLVAV	180
192	LCPKGPLRMVLTAQERNEPIFPALIYSSAAMVVTGMA-KLDPLSSQGLQLP-----	243
181	LCPKGPLRILVETAQERNEPIFPALIYSSATMVWLNFMNSAEPRNNSHVPQQENQDAV	240
243	- - - - YDPMEMEDSYDSFG - - - EPSYPEVFEPPPLTGPGEELEEEPERGVKL	286
241	APTAQPEDDGFFTPAWNQQHQHQLGPMQSTDSSREIQELPSARP- - PVEDDEERGVKL	298
287	GLGDFIEFSVLYGKAATGSDGDNNTLACFVALIGCITLILLAVERKALPALISITF	346
299	GLGDFIEFSVLYGKAATGSDGDNNTLACFVALIGCITLILLAVERKALPALISITF	358
347	GLIFYFSTDNLVRPMDFLASHOLYI	372
359	GLVFYFATDNLVYRPMFDLAVHQFYI	384

RESULT	5				
096340	096340	PRELIMINARY:	PRT;	272 AA.	
AC	096340;				
AC	096340;				
DDT	01-MAY-1999	(TREMBLEl. 10, Created)			
DT	01-MAY-1999	(TREMBLEl. 10, Last sequence update)			
DI	01-MAY-1999	(TREMBLEl. 10, Last annotation update)			
DE					
PRESENTIN	(FRAGMENT).				
DS	Drosophila melanogaster	(Fruit fly).			
OC	Eukaryota; Metazoa; Arthropoda; Insecta;				
OC	Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;				
OC	Ephydriodea; Drosophilidae; Drosophila.				

SEQUENCE FROM N.A.					
STRAIN=CANTON S.,					
MEDLINE: 98311525.					
MARFANY G., DEL-FAVERO J., VALERO R., DE JONGHE C., WOODROW S.,					
HENDRIKS L., VAN BROECKHOVEN C., GONZALEZ-DIAZ R.;					
"Identification of a Drosophila homologue: evidence of alternative spliced forms";					
J. Neurogenet. 12:41-54(1998).					
EMBL; AF017025; AAD01611.1; .					
NON TER 1	1				
SEQUENCE 272 AA:	29456 MW:	606B9A5C	CRC32;		
Query Match 30.0%; Score 576.5; DB 5; Length 272;					
Best Local Similarity 47.8%; Pred. No. 6.8e-35;					
Matches 133; Conservative 29; Mismatches 41; Indels 75; Gaps					
164 ALVFVTKLPEWSAWVILGAISYYDVLAVICPKGPLRMLVETAQERNPIFPALIYSSAMY 223					
1 ALVFVTKLPEWIAVLAISIMNDLAVISSLPRGPRLVETAQENEQIFPALIYSSTVV 60					
224 WTY-- ----- GMAKLDPSS----- QGALQLPYDPMEEDSYDSFGEDSYPE- 263					
61 YALVNTYTPQOSQATASSPSSSNSTTTTRATONSLA---SPEAAASGORTGN-SHPHQ 116					
263 ----- VFEPLTGYPGE -- ELEE----- 279					
117 NQRDDGSVLATEREAAGTQEWNSANLSVARROIEVQSTQSNGNORSNEYRTVTAPDQN 176					
279 -----EEERGVKLGLGDFTEYSVLGKAATGSGDMWNTLACEVALLIGCLTLILLAVEK 334					

Query	Match	Score	DB	Length
Best Local Matches	Similarity 73.5%;	Score 20.3%;	DB 4;	Length 184;
Matches 83;	Conservative 9;	Pred. No. 2.1e-22;		
	Mismatches 16;	Indels 5;	Gaps	
Qy	1 EELTLYGAKHVTIMLFVPTVLCKMIVVVATIKSVRFETYKEKGOLIYTPTFEDTPSYGQRLL.	60		
Dy	71 EELTLYGAKHVTIMLFVPTVLCKMIVVVATIKSVRFETYKEKGOLIYTPTFEDTPSYGQRLL.	130		
Qy	61 NSVLNTLIMISVIVMTIFLVLYKXRYCKFEHGMILMSMLFLFTIYLG.	113		
Dy	131 HSTLAAAATMSVIVMTIFLVLYKXRYCK-----VSMRHSLSLSTLFLWLG.	178		
RESULT	7			
Q19737	PRELIMINARY;	PRT;	406 AA.	
ID	Q19737; Q22692;			
AC	Q19737; Q22692;			
DT	01-NOV-1996 (TREMBLrel. 01, Created)			
DT	01-MAY-1999 (TREMBLrel. 10, Last sequence update)			
DT	01-NOV-1999 (TREMBLrel. 12, Last annotation update)			
DE	F2ZE10_5. PROTEIN.			
GN	F2ZE10_5.			
OS	Caenorhabditis elegans.			
OC	Eukaryote; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida; Rhabditina; Rhabditoidae; Rhabditidae; Peioderinae; Caenorhabditis.			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RA	GARDNER A.			
RL	Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.			
EMBL	267982; CAA9104.1;			
DR	250197; CAA9104.1; JOINED.			
EMBL	250797; CAA90577.1;			
DR	267982; CAA90677.1; JOINED.			
DR	PROSTIE; PS00319; CDP-ALCOHOL_P-TRANSF; 1.			
SQ	SEQUENCE 406 AA; 45628 MW; 07336492 CRC32;			
Query	Match 5.9%;	Score 113.5;	DB 5;	Length 406;
Best Local Matches	Similarity 24.7%;	Pred. No. 0.54;		
	Conservative 31;	Mismatches 71;	Indels 93;	Gaps
Qy	7 YGAKHVTIMLFVPTVLCKM-----IVVATIKSVRFY-----TEKGQLITYPF-T	49		
Dy	136 HGCDSTMVQFVTLNCIYAMSILGTVFYGLIVSIVSMVYCAHMSTYCGOLRSKFDVT	195		
Qy	50 EDTPSYGQRLL-----NSVLNTL-----MISVIVMTIFLVLYKXRYCKFIHGWL	97		

Qy	342 ISITFGLIFYFSTDNLVRPEMDTLASH	368	AC Q9ZC9; 1999 (TREMBrel. 10, Created)
Db	288 LSQT--LFWFLVNLF-ILKWLGSN	309	DT 01-MAY-1999 (TREMBrel. 10, Last sequence update)
RESULT	10		DT 01-MAY-1999 (TREMBrel. 10, Last annotation update)
Q35425	PRELIMINARY;	PRT;	381 AA.
ID	O35425;		
AC	Q35425;		
DT	01-NOV-1996 (TREMBrel. 01, Created)		
DT	01-MAY-1999 (TREMBrel. 10, Last sequence update)		
DT	01-MAY-1999 (TREMBrel. 10, Last annotation update)		
CYTOCHROME B.			
OS	Phascolosorex dorsalis.		"Higher-level phylogeny of Trogoniformes."
OG	Mitochondrion.		Submitted (FEB-1997) to the EMBL/GenBank/DBJ databases.
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Archosauria; Aves.		DR U81189; AAD00684; 1.
RN			KW Mitochondrion
SEQUENCE	381 AA;		SEQUENCE 380 AA; 42348 MW; 2500441E CRC32;
RP	FROM N.A.		
MLINE	93096825		Query Match 5.68; Score 108.5; DB 8; Length 380;
RA	KRAJEWSKI C., DRISKELL A.C., BAVERSTOCK P.R., BRAUN M.J.;		Best Local Similarity 20.8%; Pred. No. 1.2;
RT	"Phylogenetic relationships of the thylacine (Mammalia: Thylacinidae) among dasyuroid marsupials: evidence from cytochrome b DNA sequences."		Matches 79; Conservative 57; Mismatches 117; Gaps 20;
RT	R. Soc. Lond., B, Biol. Sci. 250:19-22 (1992).		Indels 127; Gaps 20;
CC	-1- CATALYTIC ACTIVITY: QH(2) + 2 FERICYTOCHROME C = Q + 2 FEROCYTOCHROME C.		
CC	-1- COFACTOR: TWO HEME GROUPS (B562 AND B566) WHICH ARE NOT COVALENTLY BOUND TO THE PROTEIN (BY SIMILARITY).		
CC	EMBL: M9462; AAB1692.1; -.		
DR	PFAM: PF0033; cytochrome b; N; 1.		
DR	Mitochondrion; Electron transport; Respiratory chain; Transmembrane Heme.		
KW	SEQUENCE 381 AA; 42813 MW; C6A2674F CRC32;		
SQ			
Query Match 5.7%; Score 110.5; DB 8; Length 381;			
Best Local Similarity 21.18%; Pred. No. 0.84; Indels 133; Gaps 18;			
Matches 81; Conservative 54; Mismatches 115; Gaps 18;			
Db	21 LCMIVVVATIKSYRFYTEKNGQLIYTPFEDPSVGQLLNVLNTLIMISVIVMTIFL 80		Qy 21 LCMIVVVATIKSYRFYTEKNGQLIYTPFEDPSVGQLLNVLNTLIMISVIVMTIFL 80
Db	39 MCIIQIQT-----GLFLAMHYSSTD-----LTAAFT 64		Db 40 ICLVVIQIAT -----GLMMATHYADT -----TLAESSV ----- 68
Qy	81 VVLYKRYCYKFIGWLIM-----SSMLLFELTYIYLGEVLKTYNAMDYPFLLTWN 134		Qy 81 VVLYKRYCYKFIGWLIM-----SSMLLFELTYIYLGEVLKTYNAMDYPFLLTWN 136
Db	65 SV--AHICRDVNQGWLLRNHLANGASMFMCLEHIGRIGIYSS-----LYKETWN 114		Db 68 -----AHTCRNQVOXGWLRINRHANGASMFMCLEHIGRIGIYSS-----WNTG 117
Qy	135 FGAVGMVYIHWKGSPVLIQOQAYLIMISALMAYTKLPWESAWVILGAISVYDVAVLCP 194		Qy 137 AVGAVCIHKGSPVLIQOQAYLIMISALMAYTKLPWESAWVILGAISVYDVAVLCP 196
Db	115 IG-----VILLVLMATAVGYVLPWQNSFWGATVNLMSAIPY 155		Db 118 -----VILLVLMATAVGYVLPWQNSFWGATVNLMSAIPY 158
Qy	195 KGPRLMVE-----TAQERNPPIFPALLYSSAMWTVGMAKL--DPS-SQ 236		Qy 197 PLRMLVETAQ---ERNEP-----IFPALIYSSAMWTVGMAKLDPSSQGAI--- 240
Db	156 IGP--TIAEWNGGGAYDKATLIRFFAHFILFIVITALIVLFLHETGSNNPGSGINP 213		Db 157 -----QDIVEWANGGFSDNPFLTRFALHFLPFLIASLAMHHTFHEGSNNPLGTSKC 216
Qy	237 GAOLPYDE-----MEEDSYDSFSEBPSTYEVFEP--PLTGPYGPBELEE 279		Qy 240 -OLPYDPEMEEDSYDSFG-----EPSY--PEVFEP--PLTGPYGPBELEE 279
Db	214 NAOKIPFHPPYTIKDAGFMILLISVLVLLTFPDLSGD--FDNFNSPANPLTPPHIKP 270		Db 217 DKTFPHFYPS--SLKDALGFMAMLFLLTIALFSPNLPDPHFKP-- 273
Qy	277 EEEERGGYKLGLGDFIFYSVLYGKAATAATGSGWNFTLACFLAILGLCLTLLAVFKALP 335		Qy 280 EERGKLGDFDFIFYSVLYGKAATAATGSGWNFTLACFLAILGLCLTLLAVFKALP 338
Db	271 E-----WYFLFAYAIIRSIPNKLLGG----VLAALAASVVLFLMPILHMSK-QRANT 317		Db 273 -----WYFLFAYAIIRSIPNKLLGG----VLAALAASVVLFLMPILHMSK-QRANT 317
Qy	336 ALPALPISITFGFLIFYFSTDNLV 358		Qy 339 ALPISITFGFLIFYFSTDNLV 358
Db	314 SMMFRPIQST---LFWLWLTANLJ 333		Db 318 FRPLS--QFLFWLWLTANLJ 334
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Q9ZC9	PRELIMINARY;	PRT;	748 AA.
ID	Q9ZC9; 1999 (TREMBrel. 10, Created)		
AC	Q9ZC9; 1999 (TREMBrel. 10, Last sequence update)		
DT	01-MAY-1999 (TREMBrel. 10, Last annotation update)		
DE	PUTATIVE INTEGRAL MEMBRANE EXPORT PROTEIN.		
GN	SC89.14.		
OS	Streptomyces coelicolor.		
OC	Bacteria; Firmicutes; Actinobacteria; Streptomycetaceae; Streptomyces.		
RN	SEQUENCE FROM N.A.		
RC	STRAIN=A3(2);		
RA	HARRIS D.; MURPHY L.		
RL	Submitted (FEB-1999) to the EMBL/GenBank/DBJ databases.		
RN	[2]		
RP	SEQUENCE FROM N.A.		
RC	STRAIN=A3(2);		
RESULT	11		
Q9ZC9	PRELIMINARY;	PRT;	380 AA.
ID	Q9ZC9		

Search completed: March 18, 2000, 22:07:45
 Job time: 125 sec

Query Match	Score	Length	DB	Indels	Gaps
Best Local Similarity	5.5%	382			
Matches	20.5%		Pred. No.	1.6	
Matches	79; Conservative		Mismatches	53;	
Matches	79;		Indels	139;	
Qy	21 LCMIVVVATIKSVRFTYTKNGOLIYTPTFEDTPSVQRLLNSVLNTLIMISVYMTIFL 80				
Ddb	39 MCLIQILN-----GLFLAMHXTSDT-----				-LTAFS 64
Qy	81 VVLYKRYCYKEFHGWLM-----SSLMLLEFLFTIYLGEVLTKTNVAMDYPFLTLTWN 134				
Ddb	65 SV--AHICRDVNQGWLIRNTHANGASMFNCFLHVGRGTYGST-----LYKETWN 114				
Qy	135 FGAVGMYC1HWKGPLVLQAYLIMISALMALVF1KYLPESSAW1LGALSVDLYAVLCP 194				
Ddb	115 IG-----VILLTMATATAFGYVLPQGMSFWGATVITNLSAIPI 155				
Qy	195 KGPLRMLVE-----TAQERNEPFPLIYSSAMVNTVGMAK-----LDP 233				
Ddb	156 IG--NTLIEWINGGFSVTDKATLTRFFAHPFLPFLAMVVHLFLHEIGSNNPTGLDP 213				
Qy	234 SSQGALQLQPYDP-----EMEDDSDFGEPSYPSYFEP - PLTGYPG 273				
Ddb	214 NSD---KIPFHPPYYTIKIDILGLFLMIILSLAMESPDLIGD--PDNEFTPANPLNTPEPH 267				
Qy	274 EELLEEEERVGVKLGDFIF-YSVLVGKAATGSSDWNTLACPVAILGLCLTLLAV 332				
Ddb	268 IKPE-----WYFPEFAYAIIRLSIPNKLGG --- VLLALLSILILIMPLHTST 311				
Qy	333 FKKALPALPISITFGLFYFSTDNLV 358				
Ddb	312 -QRSMMMFRPISQT--LFWMLTANLI 333				

RESULT 15

ID	PRELIMINARY:	PRT:	AA:
O35294	O35294; O35294; O35294;	PRT;	318 AA.
AC	01-JAN-1998 (TREMBrel. 05, Created)		
DT	01-JAN-1998 (TREMBrel. 05, Last sequence update)		
DT	01-MAY-1999 (TREMBrel. 10, Last annotation update)		
IDE	INTEGRIN-ASSOCIATED PROTEIN FORM 4.		
GN	TAP		
OS	Rattus norvegicus (Rat).		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;		
OC	Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.		
RN	[1]		
SRP	SEQUENCE FROM N.A.		
STRAIN	=SPRAGUE-DANLEY; TISSUE=BRAIN;		
STRA	HUANG A.M., LEE E.H.Y.;		
STRA	Submitted (AUG-1997) to the EMBL/GenBank/DDBJ databases.		
DR	EMBL: AF017437; AACB70273.1; -.		
KW	Integrin.		
SQ	SEQUENCE 318 AA; 34776 MW; FAA078CB CRC32;		

Query Match Score 106.5; DB 8; Length 318;
 Best Local Similarity 23.0%; Pred. No. 1.3; Length 318;
 Matches 40; Conservative 35; Mismatches 50; Indels 49; Gaps 8;

Qy	3 LTIKYGAH----VIMLFVPVPTLCMINIVVATIKSVRFTYTKNGOLIYTPTEDTPSVGQ 57
Ddb	158 LTIKYKSHTNKRIILLYVAGLALTIVV-----GAIIFIP-----GE 196
Qy	58 RLLNSVLNTLIMISVIVVTTFLVVLKRYCCKFHGLWIMSSMLFLFTTYLGEVLK 117
Ddb	197 KPVKNASG---LGLIVITGILLI-QINVENTAEG --MTSILVLLITQV LGYVLA 247
Qy	118 TYNVAMDPTLTLTWNFGAVGMYC1HWKGPLVLCQAYLIMISALMAYFVKYL 171
Ddb	248 VVGMCCLC1-----MACPEVGPPLLISGLGITALAEULGYMKFV 287

Om of: US-08-509-359B-137 to: GenEmbl: * out_format : pfs
 Date: Mar 18, 2000 2:37 PM

About: Results were produced by the GenCore software, version 4.5,
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Command line parameters:
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Search information block:
 Query: US-08-509-359B-137
 Query length: 448
 Database: GenEmbl: *
 Database sequences: 821193
 Database length: -1518192014
 Search time (sec): 533.740000

Search results:
 Score: 34.199999999999996

score_list:	Strd	Orig	Zscore	Escore	Len	Documentation
Sequence	gb_pr:AR060156	+ 2336.00	3900.98	3.0e-209	2285	Sequence 136 from patent US-08-509-359B-137 x AR060156
gb_pr:3:HOM51P	+ 2336.00	3900.98	2.9e-209	2229	1. Homo sapiens S182 protein (clone F-1)	
gb_pr:2:HOMSTM2R	+ 2348.00	3887.50	1.6e-208	2236	1.43964 Homo sapiens (clone F-1)	
gb_pr:AF038937	+ 2222.50	3778.58	1.9e-202	2004	1.9e-202	
gb_pr:MMU57324	+ 2232.00	3772.60	1.3e-199	1954	1.053244 Mus musculus Pg-2 mRNA	
gb_pr:AF038935	+ 2228.50	3724.54	2.0e-199	1490	1.0308935 Mus musculus presenilin 1 mRNA	
gb_pr:RNX98267	+ 2227.00	3723.07	2.4e-199	1347	1.03270 Rattus norvegicus cDNA for presenilin 1	
gb_pr:2:MMPRESEN2	+ 2224.00	3713.47	8.1e-199	2088	1.099267 R. norvegicus mRNA for presenilin 1	
gb_pr:AF0389349	+ 2218.00	3708.04	1.6e-198	1335	1.01040 M. musculus mRNA for presenilin 1	
gb_pr:AB004454	+ 2213.00	3699.29	3.9e-198	1795	1.02944 Human seven trans-membrane G protein alpha subunit mRNA	
gb Lov:D44428	+ 1624.50	204.30	1.3e-142	2490	1.00404454 Rattus norvegicus mRNA for presenilin 1	
gb_pr:AF038517	+ 1458.00	2443.57	6.4e-128	2681	1.048428 Xenopus laevis mRNA for presenilin 1	
gb_pr:AR060155	+ 1458.00	2443.82	4.2e-128	1962	1.06355 Sequence 10 from Patent AR060155	
gb Lov:MOSS1PR	+ 1468.00	2443.81	4.3e-128	1964	1.040349 Human presenilin 1 mRNA	
gb_pr:AF0389354	+ 1457.00	3701.47	8.4e-128	2791	1.04570 Sequence 133 from patent AR060154	
gb_pr:2:HUMS182R	+ 1453.50	2422.91	5.1e-128	2765	1.042110 Homo sapiens (clone CC-182) mRNA	
gb_sts:G27112	+ 1467.00	2438.57	8.3e-128	2765	1.027112 Human STS SHGC-31609, sG27112 human STS SHGC-31609, s	
gb_pr:AF06024	+ 1461.00	2428.39	3.1e-127	2791	1.063024 Sequence 1 from Patent AR06024	
gb Lov:RATS182	+ 1459.50	2433.00	1.7e-127	1407	1.0459.50 Rat mRNA for S182 protein (clone CC-182)	
gb Lov:D84427	+ 1554.50	2418.17	1.1e-126	2607	1.048427 Xenopus laevis mRNA for presenilin 1	
gb_pr:US0379	+ 1454.00	2423.87	5.5e-127	1392	1.040379 Human presenilin 1 mRNA	
gb Lov:D82363	+ 1453.50	2422.91	6.2e-127	1407	1.042363 Rattus norvegicus mRNA	
gb_pr:AR06026	+ 1448.00	2405.49	5.8e-126	3087	1.0406026 Sequence 5 from patent AR06026	
gb Lov:DRE132931	+ 1448.00	2405.60	5.7e-126	3056	1.04132931 Danio rerio mRNA for presenilin 1 (clone CC-132931)	
gb Lov:CCPS1	+ 1355.00	2258.71	8.7e-118	1241	1.04152 Cyprinus carpio mRNA for presenilin 1	
gb Lov:MMPSNINI	+ 1447.00	2411.92	2.6e-126	1430	1.027133 M. musculus mRNA for presenilin 1	
gb Lov:AF038936	+ 1438.50	2394.60	1.3e-100	1764	1.04038936 Bos taurus presenilin 1 mRNA	
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gb Lov:AF06025	+ 1418.00	2359.98	2.0e-123	1929	1.0406025 Sequence 3 from patent AR06025	
gb Lov:DRE132931	+ 1047.50	2339.94	2.6e-122	2428	1.04132931 Danio rerio mRNA for presenilin 1 (clone CC-132931)	
gb Lov:CCPS1	+ 1355.00	2258.71	8.7e-118	1241	1.04152 Cyprinus carpio mRNA for presenilin 1 (clone CC-132931)	
gb Lov:DMU78084	+ 1173.50	1950.08	1.3e-100	1764	1.0278084 Drosophila melanogaster mRNA for presenilin 1	
gb Lov:AF060183	+ 1166.50	1937.57	6.7e-100	1895	1.04060183 Sequence 165 from patent AR060183	
gb Lov:DMU77934	+ 1166.50	1938.06	6.3e-100	1809	1.0277934 Drosophila melanogaster mRNA for presenilin 1	
gb Lov:AF0170624	+ 1166.50	1937.58	6.7e-100	1894	1.040170624 Drosophila melanogaster mRNA for presenilin 1	
gb Lov:AF171064	+ 1043.00	1733.70	1.5e-88	1335	1.04171064 Caenorhabditis elegans mRNA for presenilin 1	
gb Lov:CEU5660	+ 1015.50	1686.56	6.4e-86	1461	1.03666 Caenorhabditis elegans mRNA for presenilin 1	
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gb Lov:AC014022	- 967.50	1562.38	5.3e-79	9597	1.04014022 Drosophila melanogaster mRNA for presenilin 1	
gb Lov:AF04184	+ 967.00	1599.06	4.8e-81	2601	1.04084184 Drosophila melanogaster mRNA for presenilin 1	
gb Lov:CELS35H12	+ 820.50	1328.50	5.6e-66	27102	1.041540 Caenorhabditis elegans mRNA for presenilin 1	
gb Lov:MMU57325	+ 774.00	1283.30	1.9e-63	1146	1.057325 Mus musculus PS-2 short mRNA for presenilin 1	

17	9thrSerLeuMetSerAlaGluSerProThrProArgSerCysGlnGlu	34
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34	1ArgGlnGlyProGluAspGlyLysAsnThrAlaGlnTrpArgSerGln	50
466	GCAGCAGGGCCCAGGGATGGAGAAATACTGCCAGTGGAAAGCCG	515
51	GlutGluAspGlyGluGluAspProAspartArgTyrValCysSerGln	67
516	GAGACGAGGAGGAGGTGGAGGACCCTGAGGAGGCTAACCTAACATG	565
67	YvalProGlyArgPproGlyLeuGluGluLeuThrLeuLysTyrG	84
566	GGTTCGCCGGGGCCAGGCCAGGCTGGAGGAGGCTGACCCCTCAATG	615
84	IyAlaLysIleValMetLeuIlePheValProAlaThrLeuCysMetIle	100
616	GAGGAAGATGGATGAGTCAGCTGTGCTGCTGCTGCTGCTGCTG	665
101	ValValAlaThrIleLysSerValArgPheTyrThrGluLysAsnG1	117
666	GTCGGTGTGGACCCACCATCAAGTCGCTGCGCTCTCACAGAGAAATGG	715
117	YglIleUletyrThrProPheThrGluAspThrProSerValGlyGlnA	134
716	ACAGCTCACTACAGGCCATTCACTGAGGACACCCCTCGTGGCCAGC	765
134	RgLeuLeuAsnSerValLeuAsnThrLeuIleMetIleSerValIleVal	150
766	GCCCTCTCAACTCCGGTGAACACCCCTCACTAGTCAGGCTCATG	815
151	ValMetThrIlePheLeuValValLeuTyrlsTyrArgCystYlSph	167
816	GTTATGACCATCTCTTGGTGTCTCTACAAGTACCGCTGTGTCATACTG	865
167	EileHisGlyTrpLeuIleMetSerSerLeuMetLeuLeuPheLeuHtr	184
866	CATCCATGGCTGGTTGATCATGTCCTCACTATGCTGCTCTCTCA	915
184	HrTyrIleLeuLysLeuLysThrTyrAsnValAlaMetAsp	200
916	CCPTATCATCCCTGCGTAAGGCTCATACATGTGCTGCTGCTG	965
201	TyrProThrLeuLeuThrValIrrPAsnPheGlyLalaValGlyMetVa	217
966	TACCCACCCCTCTGCTGACTGTGCGAACCTGGCCAGTGGCAGTGG	101
217	IcyIleHistTrpIleGlyProLeuValLeuGlyLalaValGlyMetVa	234
1016	GTGCATCCACTGGAAAGGGCCCTCTGGTGTCTCACAGGCCCTACCTCATCA	106
234	etileSerAlaLeuMetAlaLeuValPheIleLysTyrLeuProGluTrp	250
1066	TGACAGTCGCTCATGGCCCTAGGTTCACTGATCTGGCTGTG	111
251	SerAlaTrpValIleLeuGlyAlaIleSerValTyrAspLeuValAlaVa	267
1116	TCCCGTGTGGTCATCTGGGCCATCTCTGTGATCATCTGGCTGT	116
257	LeuGlyProTrpGlyProLeuArgMetLeuValGluThrAlaGlyGluA	284
1116	GCTGTGTCCCAGGGCCCTCTGAGAATGCTGTAGAAACTGCCAGGGA	121
284	RgAsnGluProIlePheProAlaLeuIleTyrSerSerAlaMetValTrp	300
11216	GAATTAGGCCATATGCCGCTCATATGCTGCTGCTGCTGCTG	126
301	ThrValGlyMetAlaLysLeuAspProSerGlnGlyAlaLeuGlnLe	317
1266	ACGGTGGCATGGCCAAAGCTGACCCCTCCCTCAAGGTGGCCCTCCAGCT	131
317	uProtTrpAspProGluGluAspSerGlyAspSerPheGlyGluP	334

1042	CCTCATACCCGAAGCCCTTCGAGCCCCCTGCCCTGCTGCTACCCAGGGAG	1091
351	GluLeuGluGluGluGluGluGluGluGluGlyvalysLysGlyLysGlyAspPh	367
1092	GAGCTGAAGGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGACTT	1141
367	ellephenylservalinevalylGlyLysAlaAlaAlaLysAlaLysAlaLysGly	384
1142	CATCTTACAGCTGTGGCAAGGCTGAGGACTGCAGAACGGAG	1191
384	SPTPAsparthThrLeuAlaCysPheValAlaIleLeuIleGlyLeuCys	400
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401	LeuThrLeuLeuLeuAlaValPhyLysLysAlaLeuProAlaLeuPr	417
1242	CTCACCTCCGCTGTGCTGTGCTGTGCTGAAGGGCTTGGCCCTCCC	1291
417	oilSerSerIleThrPheGlyLeuIlePheTerPheSerThrAspAspIleu	434
1292	CATCTCATCACCTGGACATCTACTCTACAGACAACCTGG	1341
434	aLArgProProMetAspThrLeuAlaSerHisGlnLeutYile	448
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seq_documentation_block:		
LOCUS	AF038935	1490 bp
DEFINITION	Mus musculus presenilin 2 mRNA	mRNA
ACCESSION	AF038935	complete cds.
VERSION	AF038935.1	
KEYWORDS	GI:2724097	
SOURCE	house mouse.	
ORGANISM	Mus musculus	
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; M		
Eutheria; Rodentia; Sciuromorpha; Muridae; Murinae; M		
AUTHORS	Sahara, N., Mori, H. and Shirasawa, T.	
TITLE	Molecular cloning of mouse presenilin 2 gene	
JOURNAL	Unpublished	
REFERENCE	2 (bases 1 to 1490)	
AUTHORS	Sahara, N., Mori, H. and Shirasawa, T.	
TITLE	Direct Submission	
JOURNAL	Submitted (16-DEC-1997) Molecular Biology, Tokyo Institute of Psychiatry, 2-1-8 Kami-kitazawa, Setagaya-ku, Tokyo 153-0043, Japan	
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315 a	418 c	416 g
315	418	341 t

501 CATCCATGGCTGGTGTGATCATCTCCCTCCTGATGCCTCTTCGGTCA 550
 184 hrtyriletryleGlyGlutLeuLysThrTyrAsnValAlaMetASP 200
 551 CCTACACTACCTCGGAAGTTCAAGCTACATGTGCCATRGAC 600
 201 TyrProThrLeuLeuLeuThrValTyrPAsnPheGlyAlaValGlyMetVa 217
 601 TACCCACACACTCTGGCTGTCGAACCTCGGGCAGTGCGATGGT 650
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 651 GTGGATCACACTGAAAGGTCCCTGGTGTCAAGAGCTAACCTACATGT 700
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 ACCESSION X99467
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 VERSION 1
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 SOURCE Rattus norvegicus
 ORGANISM Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
 Rodentia; Sciurognathi; Muridae; Murinae; Rattus .
 REFERENCE 1 (bases 1 to 2088)
 AUTHORS Frentzel,S., Abel, A.S. and Luebbert, H.
 JOURNAL Unpublished
 2 (bases 1 to 2088)
 AUTHORS Frentzel,S.
 TITLE Direct Submission
 JOURNAL Submitted (05-JUL-1996) S. Frentzel, Sandoz Pharma Ltd, Preclinical Research, Po Box, CH-0002 Basel, Switzerland
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 171 GACCTCCCTGATGTAGCCCTAGGCCAGAGCCCTATCACGTTCCGCCAGACA 220
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seq_documentation_block :

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 DEFINITION Human seven trans-membrane domain protein (AD3LP/AD5) mRNA,
 complete cds.

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 VERSION U34349 1
 KEYWORDS Alzheimer's disease.
 SOURCE Homo sapiens
 ORGANISM Eukaryota; Primates; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 1673)

REFERENCE Li,J., Ma,J. and Potter,H.
 AUTHORS Li,J., Ma,J. and Potter,H.
 TITLE Identification and expression analysis of a potential familial
 Alzheimer disease gene on chromosome 1 related to AD3
 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 92 (26), 12180-12184 (1995)
 MEDLINE 96109229
 FEATURES 1. 1705
 REFERENCE 2 (bases 1 to 1673)
 AUTHORS Li,J.
 TITLE Direct Submission
 JOURNAL Submitted (21-AUG-1995) Jinhe Li, Neurobiology, Harvard Medical
 School, 220 Longwood Ave., B2-502, Boston, MA 02115, USA
 LOCATION/Qualifiers 1. 1705
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ACCESSION	LA2177			
VERSION	LA2177.1	GI:904129		
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;				
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
REFERENCE	1 (bases 1 to 1964)			.Le 75
AUTHORS	Shearrington,R., Rogaev,E.I., Liang,Y., Rogaeva,E.A., Levesque,G., Ikeda,M., Chi,H., Lin,C., Li,G., Holman,K., Tsuda,T., Mar,L., Poncin,J.-F., Bruni,A.C., Montesi,M.P., Sorbi,S., Rainero,I., Pinessi,L., Nee,L., Chumakov,I., Pollen,D., Brookes,A., Sanseau,P., Polinsky,R.J., Wasco,W., Da Silva H.A.R., Haines J.L., Pericak-Vance,M.A., Tanzi,R.E., Ross,A.D., Fraser,P.B., Rommens,J.M. and St. George-Hyslop,P.H.			
TITLE	Cloning of a gene bearing missense mutations in early-onset familial Alzheimer's disease			
JOURNAL	Nature 375 (6534), 754-760 (1995)			
MEDIUM	95319502			
COMMENT	On Jul 25, 1995 this sequence version replaced gi:897616.			
FEATURES	Location/Qualifiers			

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OM of: US-08-509-359B-137 to: N_Geneseq_36:*
out_format : pfs

Date: Mar 18, 2000 11:46 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-1998 Compugen Ltd.

Command line parameters:
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-DELPOP=6.000 -DDELEXT=7.000 -SPART=1 -MATRIX=Hlossum62
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-ALIGN=15 -MODE=LOCAL -OUTFORMAT=pfs -NORMEXT -MINLEN=0
-MAXLEN=1000000 -USER=US08509359 -NCPU=6 -ICFU=3 -NO_XLPPXY -WAIT
-THREADS=1

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Query length: 448
Database: N_Geneseq_36:*
Database sequences: 311585
Database length: 125096042
Search time (sec): 37.450000

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N_Geneseq_36:T40038	+	186.00	319.87	3.2e-10	996	Early onset Alzheimer's dise
N_Geneseq_36:T40038	+	167.50	292.61	1.1e-08	540	Presenilin-1 exon 9.

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-QGAPOP=4.500 -OCAPEXT=0.000 -XGAPEXT=0.500
-EGAPOP=6.000 -IGAPENT=7.000 -YGAPOP=10.000 -YGAPEXT=0.500
-DELTRANS=6.000 -DELEXT=7.000 -START=200 -MATRIX=I6osm62
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Database sequences: 311585
Database length: 125096042
Search time (sec): 37.1500

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N_Genesis	-36..V9555	+ 1460..00	2675..97	1..9e-141	2764	1 Homo sapiens PS1 cDNA. Nucleic	
N_Genesis	-36..T59536	+ 1454..00	2670..06	4..0e-141	1762	1 Human early onset Alzheimer's	
N_Genesis	-36..T87432	+ 1438..00	2670..14	4..0e-141	1750	1 Human early onset Alzheimer's	
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N_Genesis	-36..V04667	+ 1448..00	2652..38	3..9e-140	3086	1 Human presenilin-1 cDNA (hPS1)	
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N_Genesis	-36..T63227	+ 1442..00	2646..81	7..9e-140	1911	1 Human S182 gene associated with	
N_Genesis	-36..T87432	+ 1438..00	2642..29	1..4e-139	1488	1 Partial AD3 sequence. Identify	
N_Genesis	-36..T85333	+ 1435..00	2640..50	7..1e-139	1703	1 PS1/429 protein coding sequence	
N_Genesis	-36..T40043	+ 1166..50	2134..83	2..6e-111	1895	1 Human mutant S182 gene, PS1.1c	
N_Genesis	-36..T60336	+ 1003..50	1834..59	1..4e-94	1500	1 Presenilin homologue. New pres	
N_Genesis	-36..T51217	+ 666..00	1220..72	2..1e-60	473	1 Caenorhabditis elegans SEL-12	
N_Genesis	-36..T51258	+ 385..00	679..56	3..0e-30	2387	1 Human expressed sequence tag ES	
N_Genesis	-36..T51256	+ 1435..00	2640..50	7..3e-139	2765	1 Human presenilin-1 gene genomic sequ	
N_Genesis	-36..T85333	+ 1435..00	2629..50	7..3e-139	2765	1 Human presenilin-1 gene genomic sequ	
N_Genesis	-36..V12811	+ 348..00	638..06	6..1e-28	230	1 Human biallelic polymorphic DNA	
N_Genesis	-36..X11761	+ 348..00	638..06	6..1e-28	230	1 Human biallelic polymorphic DNA	
N_Genesis	-36..T40037	+ 315..00	560..25	1..3e-23	945	1 Presenilin-1 exon 8. New presen	
N_Genesis	-36..T99636	+ 315..00	560..25	1..3e-23	945	1 Human presenilin-1 gene exon 8	
N_Genesis	-36..T51200	+ 270..00	459..79	5..2e-18	4004	1 Human AD4 gene genomic sequen	
N_Genesis	-36..T51259	+ 226..00	452..00	1..3e-17	2058	1 Human AD4 gene genomic sequen	
N_Genesis	-36..T51259	+ 229..00	396..44	1..8e-14	1438	1 Human AD4 gene genomic sequen	
N_Genesis	-36..T40052	+ 224..00	394..02	2..4e-14	736	1 Presenilin-1 exon 13. New pres	
N_Genesis	-36..V99671	+ 224..00	394..02	2..4e-14	736	1 Human presenilin-1 gene exon 13	
N_Genesis	-36..T40034	+ 212..50	362..71	1..3e-12	1727	1 Presenilin-1 exon 5. New presen	
N_Genesis	-36..T99654	+ 201..00	340..33	2..3e-11	1883	1 Presenilin-1 exon 6. New pres	
N_Genesis	-36..V99654	+ 201..00	340..33	2..3e-11	1883	1 Presenilin-1 exon 6. New pres	

N_Geneseq_36:T40041	+	191.00	329.08	9.9e-11	1003	Presenilin-1 exon 12. New P
N_Geneseq_36:T59973	+	191.00	329.08	9.9e-11	1003	Human Presenilin-1. Gene exo
N_Geneseq_36:T40038	+	186.00	319.87	3.2e-10	996	Early onset Alzheimer's dise
N_Geneseq_36:T40038	+	167.50	292.61	1.1e-08	540	Presenilin-1 exon 9.

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seq_documentation_block:
ID T40031 standard; DNA; 2229 BP.
AC T40031;
DT 25-JUL-1997 (first entry)
DE Human presenilin-2 wild type coding sequence.
PR presenilin-2; human; hPS1-2; PS-2; integral membrane protein; AD;
KW Alzheimer's disease; cerebral haemorrhage; schizophrenia;
KW depression; antibody; gene expression modulator; therapy; ss.
OS Homo sapiens.
PH Key Location/Qualifiers
      FT 366..1712
      FT /*tag= a
      FT /product- presenilin-2

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acid probes specific for the mutant gene, provides a means of diagnosing Alzheimer's disease.

Sequence 2236 BP; 488 A; 584 C; 645 G; 519 T;

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alignment_scores:
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  Ratio: 5.214          Gaps: 0
  Percent Similarity: 100.000   Percent Identity: 100.000

alignment_block:
  US-08-359B-137 x T51253 .. .
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  1 MetLeuThrPheMetAlaSerAspSerGluGluValCysAspGluAla 17
  368 ATGCCACATTCATGCCCTCTGACGGCGAGGAAGACTGTGAGGCG 417
  17 gThSerLeuMetSerAlaLysSerProThrProArgSerCysGlnGluG 34
  418 GACGCCCCATATGCGCCAGGCCAGCGGGCTCTGCCAGGAGG 467
  34 IYArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50
  468 GCAGCAGGGCCCAAGGGTGGAGAACAATGCCAGTGAGAAAGCCAG 517
  51 GluAsnGluLysAspGlyGluGluAspPTOAspArgTyValCysSerG1 67
  518 GAGAACGAGGGAGGGTGGAGGGACCCCTGACCCCTATGTCGAGTGG 567
  67 yValProGlyArgProProGlyLeuGluGluLeuThrLeuLysTyrg 84
  568 GGTTCCGGGGCCAGGCCAGGAGGAGGAGAACAATGCCAGTGAGAA 617
  84 IYAlaLysHsValLeuMetLeuPheValProValThrLeuCysMetIle 100
  618 GAGCCAAAGCAGCTGATCATGCTGTTGCTGTACTCTGTGCTATGATC 667
  101 ValValValAlaThrIleSerValArgPheThrGluLysAsnG1 117
  668 GTGGGGGTACCCACCATCAAGTCAGTCAGTCAGTCAGTCAGTCAGTC 717
  117 yGlnLeuIleIrrhProThrGluAspThrProSerValGlyGlnA 134
  718 ACAGGTCATCTACAGCCATTCAACAGGACACACCCTCAGTCAGTCAGTC 767
  134 rGLeuLeuAsnSerValLeuAsnThrLeuIleSerValIleVal 150
  768 GCCCTCTCAACTCCGGCTGTAACACCCCTCATCATACCGTCATCGTG 817
  151 ValMetThrIlePheLeuValLeuIleLeuIleLeuIleLeuIleLeu 167
  818 GTTATGACCATCTTCTGTTGCTGCTACAAGTACCGCTGCTACAGTT 867
  167 eIleIleHisGlyTrpIleIleMetSerSerIleLeuLeuLeuLeuPheI 184
  868 CATCATGCTGGTTGATCATGCTCACTGATGGTGTGTTCCCTCTCA 917
  184 hrTyryIleLeuGlyGluValLeuLysThrTyAsnValAlaMetAsp 200
  918 CCTATATCACCTGGAAAGTCCCAGACCTCATCATGTCATGGCAGGGAC 967
  201 TyrProThrIleLeuLeuIleLeuIleLeuIleLeuIleLeuIleLeu 217
  968 TACCCACCCCTCTGCTGACTGCTGAACTCGGGCAGTGGGATGGT 1017
  217 IcysIleHistDpIleGlyProLeuValLeuIleGlnAlaTyrIleIleM 234
  1018 GRCATCCACTGGAAAGGGCCCTCTGCTGCTGCACTGGGGCCTACCCATCA 1067
  234 etIleSerAlaLeuAlaLeuValPharleLysTyLeuProGluTrp 250
  ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
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1068 TGATCAGTAGCCCTCATGGCCCTAGTGTTCATCAAGTACCTCCAGACTGG 1117
 251 SerAlaItpValAlleLeuGlyAlaIleSerIleValtyAspLeuValAla 267
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 1118 TCCGCCTGGGTCATCTGGGGCATCTCTCTGTTGATCATCTGTTGGCTGT 1167
 267 ILeuCysProIysGlyProLeuArgMetLeuValGluThrAlaGlnGluA 284
 1168 GCTGTGTCGCCAAAGGSCCTCAGAAGTGTGTTAGAAACTGCCAGAGA 1217
 284 rGAsnGluProIleIlePheProAlaLeuIleSerSerAlaMetValTrp 300
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 1218 GAATGAGCCATATCCCTCGTATACTCATGCTGTCATGGCTGG 1267
 301 ThrValGlyMetAlaLysLeuAspProSerSerGlnGlyAlaLeuGlnE 317
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 1268 ACGGTTGGCATGGCAAGCTGTCAGCCCTCCCTCAGGGTGCCTCCAGCT 1317
 317 uProTyRAspProGluMetGluGluAspSerTyAspSerPhGlyGluP 334
 1318 CCCCTAGACCCGGAAATGGGAGAACTCTATGACAGTTTGGGAGC 1367
 334 roSerTyrProGluValPhGluIuProProLeuThrGlyTyProlGlyGlu 350
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 1368 CTTCATACCCGAAGCTCTGACTGCTACCCAGGGAG 1417
 351 GluIleGluGluGluGluGluGluGlyAlaIleLeuGlyAlaLeuGlyAspPh 367
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 1418 GAGCTGAGGAGGAGGAGGAGGAAAGGGCTGTGAGCTTGCCCTGGGACTT 1467
 367 eIlePheryrSerValIeUalGlyLysAlaAlaIaLathGlySerGlyA 384
 1468 CATCTCTACTGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTC 1517
 384 SPTRPAsnThrThrLeuAlaCysPheValAlaIleLeuIleLeuIleLeu 400
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 1518 ACTGGATAACACGTCAGCCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1567
 401 LeuThrLeuLeuAlaValPhLeuLysAlaIleLeuIleLeuIleLeuIle 417
 1568 CTGACCCCTCCCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1617
 417 oIleSerIleIrrhPheClyLeuIleLeuIleLeuIleLeuIleLeuIle 434
 1618 CATCTCCATCTACGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTCAGTC 1667
 434 aIArgProPheMetAspThrIleAlaSerHisGlnLeuTyrtle 448
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 1668 TCGGGCGGTTATGGACACCTGGCCCTCCATCAGCTCATC 1711

seq_name: N_Geneseq_36:v04669

seq_documentation_block:

- ID V04669 standard; cDNA: 2229 BP.
- AC V04669
- DT 20-JUL-1998 (first entry)
- DB Human Presenilin-2 cDNA (hPS2).
- KW Presenilin-1; PS1 gene; human; familial Alzheimer's disease; FAD; cerebral haemorrhage; schizophrenia; depression; epilepsy; mental retardation; diagnosis; therapy; transgenic animal; ss. Homo sapiens.
- OS Homo sapiens.
- Key Key
- PH CDS
- FT /*tag= a
- FT mutation
- FT /*tag= P
- FT mutation
- FT /note= "A to T FAD mutation site (Asn141le)"
- FT 1080
- FT /note= "T to G FAD mutation site (Met239Val)"
- FT 1624
- FT /note= "T to C FAD mutation site (ile420Thr)"
- FT

344	GGCGGCCAAGTAGTAACGTTGAAACAGATGAGGAGGA	393
seq_documentation_block:		
ID T40030 standard; DNA: 1964 BP.		
T40030;		
DT 23-JUL-1997 (first entry)		
Murine presenilin-1 wild type coding sequence.		
Presenilin-1; mouse; hps1-1; hps1-2; PS-2; integral membrane protein; AD;		
familial Alzheimer's disease; cerebral haemorrhage; schizophrenia;		
depression; antibody; gene expression modulator; therapy; ss.		
Mus musculus.		
Key		
LocationQualifiers		
cds	188..1951	
/*tag= a		
/product= presenilin-1		
FT W0934099-A2.		
PN PD 31-OCT-1996.	CA0263	
PF 29-APR-1996.	CA0263	
PR 28-APR-1995; US-431048.		
PR 28-JUN-1995; US-496841.		
PR 31-JUL-1995; US-509359.		
(HSCR-) HSC RBS & DEV LP		
(PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.		
PA Fraser PE, Rommens JM, St George-Hyslop PH;		
WPI; 96-497631/49.		
P-PDB; W05735.		
DR		
New presenilin genes - useful for diagnosis, therapy and drug screening of familial Alzheimer's disease, cerebral disorders, etc.		
PT This sequence represents the coding sequence for the murine presenilin 1.		
CC Claim 8; Page 145-146; 178pp; English.		
CC This sequence represents the coding sequence for the two different forms of wild type human presenilin-1 (PS-1). The form represented by T40029 results from alternate splicing of the genomic DNA sequence.		
CC T40031 represents the coding sequence for wild type human PS-2. The presenilins are a family of highly conserved integral membrane proteins with a common structural motif, common alternate splicing patterns, and common mutational hot spot regions. Mutations in PS genes are implicated in familial Alzheimer's disease (AD) and possibly other diseases such as cerebral haemorrhage, schizophrenia, depression etc., so detection of mutations in these sequences can be used for diagnosis of these diseases.		
CC The encoded proteins, or vectors that express them or containing antisense sequences, antibodies selective for mutant forms of the encoded proteins (such as W05736) and modulators of PS gene expression are potentially useful for treatment of AD etc. Transgenic animals are useful as models for drug screening. The antibodies can also be used e.g. for affinity purification and in immunassays.		
SQ Sequence 1984 BP; 503 C; 496 G; 460 T;		
alignment_scores:		
Quality: 1468.00	Length: 467	
Ratio: 3.915	Gaps: 6	
Percent Similarity: 80.300	Percent Identity: 64.026	
alignment_block:		
US-08-359B-137 x T40030 ..		
Align seg 1/1 to: T40030 from: 1 to: 1964		
seg_documentation_block:		
Quality: 1468.00	Length: 467	
Ratio: 3.915	Gaps: 6	
Percent Similarity: 80.300	Percent Identity: 64.026	
seg 24 GluUserProThrProArgSerCysGlnGluGlyArgGlnGlyProGluAs 40		
194 GAGATACCTGCACCTTGTCCTACTTCAGGATGCCAGATGTCGAGGA 243		
40 GlyGluAspAspAspAspAspAspAspAspAspAspAspAspAspAspAsp 57		
244 CAGGCACTGCCAGCAGGCCATCCGGACGAGACAGAACGACAGTCGTTCTGGGTCTGGGACGTC 293		
57 LysGluAspAspArgTyryValCysSerGlyValPro..... 69		
294 AGCAAGCAGCATGACAGGCAGAGACTGACAACTCTGAGCCAAATCTAAAT 343		
50 GlyLysProGly.....Le 75		

292 euIletryrSerAlaMetValTrpThrValGlyMetAlaLysLeuAsP 308
 1111 TTATCATTCTCAACAATGGTGTGAAATGGCTGAAGGAGAC 1160
 DR p-PSDB; W23966.
 PT New isolated mutant presenilin-1 genes - useful for developing
 PT products for use in detection, diagnosis and therapy of Alzheimer's
 PT disease and for drug screening
 PS Disclosure; Page 197-199; 238pp; English.
 CC This cDNA clone for a murine presenilin-1 (PS1) homologue (see
 CC W23966). It was isolated from a mouse cDNA library using a DNA
 CC probe from the human presenilin-1 (PS1) gene (see V04666). Mutations in the
 CC human PS1 and PS2 genes (see V04666-68) have been linked to the
 CC development in humans of forms of familial Alzheimer's disease
 CC (FAD). All amino acids that are mutated in analysed FAD
 CC pedigrees were conserved in the murine homologue. Use of the
 CC nucleic acids and proteins comprising or derived from presenilins
 CC can be made in screening and diagnosing FAD, identifying and
 CC developing therapeutics for treatment of FAD, and in producing cell
 CC lines and transgenic animals useful as models of FAD. Sequence 1964 BP;
 SQ 503 A; 496 G; 460 T;

309 ProSerSerGlnGlyAlaLeuGlnLeuProTyroProGluMetGlu.. 324
 1161 CCAGAAGCCCCAA.....AGAGGGTACCCCAAAGAACCTAACAC 1204
 325GlutAspSerTyrAspSerPheGlyGluP 334
 1205 ACAAAAGCGGAGAGAGAGACAGGACAGGGTGTGGAAACGGATG 1254
 334 roSerTyrProGluValAlpHeGluProProLeuThrGlyTyroGly... 349
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 1305 CATGGCTCCACTCCCACTGTCAGAGGTGCTCAGGAACATTCGGAG 1354
 353Glugluleu..... 365
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 365 LyAspPheIlePheTyrSerValLeuValGlyLysAlaAlaAlaAlaArgly 381
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 382 SerGlyAspTyrPheSerThrIleLeuAlaCysPheValAlaLeuIleG1 398
 1455 AGTGGGAGACTGGAAACCAACATAGCTGCTTGTAGCCATACTGATCGG 1504
 398 YLeuCysLeuThrIleLeuLeuAlaValA.PheTysLysAlaLeuPro 415
 1505 CCTGTCCCTAACATACTCCCTGCTCCCAATTTCAGAAACGGTCCAG 1554
 415 IaLeuProIleSerIleThrPheGlyLeuIlePheTyrPheSerThrAsP 431
 1555 CCCTCCCACATCCCATCACCTTGCGCTCGCTGTTTACTGGCCAGGGAT 1604
 432 AsnLeuValArgProPheMetAspThrLeuLeuSerHisGlnLeuTyrI1 448
 1605 TACCTGTGAGCCCTCATGGACCAACTIGCATCCATCAGTTATAT 1654
 448 e 448
 1655 C 1655

seq_name: N_Geneseq_36:V04666
 seq_documentation_block:
 ID V04666 standard; cDNA: 1964 BP.
 AC V04666;
 DT 20-JUL-1998 (first entry)
 DE Mouse Presenilin-1 cDNA.
 KW presenilin-1; PS1 gene; mouse; familial Alzheimer's disease; FAD;
 KW cerebral haemorrhage; schizophrenia; depression; epilepsy; ss.
 KW mental retardation; diagnosis; therapy; transgenic animal; ss.
 OS Mus musculus.
 FH Location/Qualifiers
 FT 188 .1561
 FT /*tag= a
 PN W09801549-A2.
 PD 15-JAN-1998.
 PF 04-JUL-1997; CA0475.
 PR 02-JAN-1997; US-034590.
 PR 05-JUL-1996; US-024673.
 PR 12-JUL-1996; US-024700.
 PR 08-NOV-1996; US-024995.
 PA (HSCR-) HSC RES & DEV LP.
 PA (UTOR) UNT TORONTO GOVERNING COUNCIL.
 PI Fraser PE, Rommens JM, St George-Hyslop PH;
 DR WPI: 98-286355/25.

75 .LeuGluGluGluLeuThrIleUlyStryGlyAlaLysHisValIleMetL 91
 452 AGAACATGGAGCTGACAUTGAATAATGCCAAGCAATGTGACATGC 501

91 eUphevalProvalThrLeucystmetilevalvalAlaIthrIleLys 107
 502 TCTTGTCCCTGTGACTCTGATGGTGGCTACCTTAAG 551

108 SerValArgPheYrThrGluLysAsnGlyGlnIleLeuIleYrThrProPh 124
 552 TCAGCAGCUTTTAACCGGAAGGATGGCAGCAATTATAACCCATT 601

124 eThrGluAspThrProSerValGlyGlnArgLeuLeuAsnSerValLeuA 141
 602 CACAGAAGATAACCCGAGACRTGGGCCAGAGGCCCTGCACTCAATCTGA 651

141 snthLeuIleMetIleSerValIleLeuIleMethIleLeuIleLeuIle 157
 652 ATGCTGCCATCATGTCAGTCAGTGTCACTACTATCCTCTGGTG 701

158 ValLeutYrLysTrpArgCystYrLysPheIleHisGlyTriPheIleMe 174
 702 GTCUGTATAATAGGTCTATAAGTCATCCATGCCCTGGCTTATTAT 751

174 tSerSerLeuMetIleLeuPheLeuPheThrIleYrLeuGlyGluV 191
 752 ATCACATCPATTGTTGCTGTTCTTTTCTTCTTCTTACTTCATTCTTAAAG 801

191 alleLysThrItyrAsnValAlaMetAspTyPthrIleLeuLeuIthr 207
 802 TGTAAAACCTATAACGTGACTACATGTTGACTGACTGCACTCCRG 851

208 ValTrpAsnPheGlyAlaValGlyMetValCysIleHistPlysGlyPr 224
 852 ATCAGGAATTGGTGGTGGGATGATTCATTCACCTGGAAAGGTCC 901

224 oLeuValLeuGlnGlnAlaItyrLeuIleMetIleSerAlaLeuMetAlaL 241
 902 ACGTGACCTCCAGGAGGCAATCTCATGGCTTCATGGCCC 951

241 euvalPheIleItyrLeuProGlyItrpSerAlaItrpValIleLeuGly 257
 952 TGGTTTATCAAGTACCTCCCTGATGGCTGCTGGCTCATCTGGCT 1001.

258 AlaIleSerValYrAspIleuAlaValLeuCysProIysGlyProle 274
 1002 GTGATTTCAGTATATGATTAGTCGCTGTTGTCAGAAAGTCCACT 1051.

274 uATGmetLeuValGluThrAlaIgluIuaGasnGluProIlePheProA 291
 1052 TCGPATGCTGGTTCAAACAGCTCAGAGAAATGAAACCTTTCAG 1101.

291 IaleuIleYrSerAlaMetValItrpIhrGlyMetAlaLysLeu 307
 1102 CTCTATTAATCTCTAACATGGTGGTGGAAATGGCAAAAGGA 1151.

308 AspProSerSerGlnGlyAlaLeu GlnLeuProTyAspProG1 322
 1112 GACCGGAAGCTCAAGGAGATCCAAGAAATTCCAAGTATAATGCAAGA 1201.

322 u....MetGluGluAspSertyrasSerPheGlyGlu.....ProS 335
 1202 AAGCACAGAAAGGGGATCCAAGCAAGACTGTCAGAATGATGGCCG 1251.

335 ertryProGluvalPheGluProLeuIhrGlyTyProlG1 349
 1252 GGTCAGTGAGGAATGGAAAGCCAGGGACAGCATACTAGGGCCTCAT 1301.

350GluGluIle 352
 1302 CGCTCATACCTGACTCAGGACTTCAGGACTTCAGGAGTAT 1351.

353GluGluGluGluGluGlyIvalIysLeuGlyA 366

1352 CCTCGCTGGTAAAGACCAGGAAAGGGAGAAAACCTGGATTGGGAG 1401
 366 sPheIIlephenryrSerValLeuValGlyLysAlaAlaIalathGlySer 382
 1402 ATTCAATTCTCAACAGTGTTCAGGCTAGCAAGGCCAGT 1451

383 GlyAspItpAsnThrThrLeuAlaCysPheValAlaIleLeuIleGlye 399
 1452 GGACGGAACAACATGGCTGTTGACATATAATGGTT 1501

399 uCysLeuThrIleLeuLeuAlaValPhelysLysAlaLeuProAlA 416
 1502 GTGCCCTAACATTAACTCCCTGCCCCATTTCAGAAACCATTCAGCTC 1551

416 eUprolleserIleRlIlePhGlyLeuIlePhenryrPheserThrasphn 432
 1552 TICCAAACTCTCATCACCTTGGCTTGTTCTPACTTNGCAGAGATAT 1601

433 LeuValArgProPheMetAspItpLeuAlaSerIleuAsnLeytrylle 448
 1602 CTTGTAAGGCCTTTAGGACCAATTAGCATCAATTCATCAATTTAC 1649

seq_name: N_Geneseq_36:v17358

seq_documentation_block:
 ID V17358 standard; DNA; 2764 BP.
 AC V17358;
 DT 04-JUN-1998 (first entry)
 DE PS1/467, protein coding sequence.
 KW Presenilin Peptide; PS1/429; ImmunoGen; immune response; PS1 gene;
 KW Alzheimer's disease; mitochondrial pathology; neurodegeneration;
 KW apoptosis; PS1/467; ss.
 OS Homo sapiens.
 FH Location/Qualifiers
 Key 249 . 1652
 CDS FT
 FT PN W09746678-A1.
 FT PD 11-DEC-1997.
 PS PF 03-JUN-1997; US09272.
 CC PR 06-JUN-1996; US-6833315.
 CC PA (FARB) BAYER CORP.
 PI Chisholm JC, Davis JN, Drache B;
 DR WPI; 98-042186/04.
 DR P-PSDB; W4430.
 PT DNA encoding presenilin peptide PS1/429 and its analogues - useful
 for diagnosis and treatment of Alzheimer's disease
 Disclosure: Fig 2; 77pp; English.
 This sequence encodes the PS1/467 presenilin peptide. This sequence is
 specifically stated as not being in the nucleic acid of the invention,
 which encodes the PS1/429 presenilin peptide PS1/429 (II). Cells
 transformed with the DNA are used to produce recombinant (II) and
 analogues, useful e.g. as immunogens for generating an immune response
 against PS1/429. (II) is a new product of the PS1 gene, mutations in
 which cause Alzheimer's disease (AD). The nucleic acids are generally
 used as probes for detection and quantification of PS1/429,
 particularly for diagnosis of AD, especially the target sequences that
 hybridise with probes are isolated for sequencing. Antibodies (Ab) can
 also be diagnosed at the protein level using Ab as immunoassay reagents.
 Ab can also be used to identify epitopes and for affinity purification of
 peptides. Antisense nucleic acid may also be used to regulate expression
 of the PS1/429 gene, and both nucleic acids and peptides are useful as
 size markers in electrophoresis, chromatography etc. The transgenic
 animals are used as models for AD, e.g. for testing drugs. Regulators of
 the PS1/429 gene or polypeptide can be used to treat e.g. AD or diseases
 involving mitochondrial pathology, apoptosis and neurodegeneration.
 Typical regulators are antisense sequences, ribozymes, aptamers,
 synthetic or natural compounds. (II) may also be used to target other
 coding sequences to particular cellular locations.
 SQ Sequence 2764 BP; 715 A; 624 C; 653 G; 772 T;

Antibodies against the mutant polypeptide can also be used for this purpose. Vectors containing or expressing a nucleic acid molecule, protein or antibody specific for mutant PS1 can be administered to a patient to reduce the likelihood, or delay the onset, of Alzheimer's disease, e.g., anti-sense RNA expression can be used to decrease expression of the PS1 peptide. Transgenic animals expressing the Alzheimer's disease protein can be used to test candidate therapeutics and to investigate the normal role of PS1. The PS1 peptide may also be included in pharmaceutical compositions (vaccines) for Alzheimer's disease therapy.

852	ATCTGGAATTGGTGTGGATGTTCCATTACTGGAAAGGCC	901
224	oleuvalleuGlnAlaItylLeuIleMetIleSerAlaLeuMetAlaL	241
902	ACITGACTCCAGCAGCATATCCTATTGATGATGTCATCTGGCCC	951
241	euaPheIleIlystYtrIleProGluTrpSerAlaTrpValIleugly	257
952	TGGTTTATCAAGTACCTCCCTGAATGACTGGTGGTCATCTGGCT	1001
258	AlaIleSerValItyAspIleuValAlaLeucySProIysGlyProte	274
11002	GGATGTTCACTATAATGATTAGTTAGTGCGTGTGTCAGCTAAC	1051
274	wargMetLeuValGluThrAlaIglnGluArgAsnGluProIleHePro	291
11052	TCGTATGCTGGTGAACAGCTCAGAGAAGAAATGAAACGCTTTCCAG	1101
291	IleauIleIlyserSerAlaMetValTrpIrrhYalGluMetAlaLysLeu	307
11102	CCTCTATTACTCTCAACATGGTGTGTTGGTGAATATGGCAAGAGGA	1151
308	AspProSerSerGlnGlyAlaLeu.....GlnLeuProTyraSProG1	322
11152	GACCCGGAGCTAAAGGAGATCCAAAATTCACCACTAAATGCAGA	1201
322	u...MetgluGluAspSerTyraPheGlu.....ProS	335
11202	AAGCACAGAAAGGAGTCACAAAGACACTGTCAGAGAATGATGATGGGG	1251
335	erytYProGluValPheGluProProLeuThrGlyTyPheGly.....	349
11252	GGTCAGTGAGGATGGAGCCAGAGGACAGTCATCTGGCCTCAT	1301
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11302	CGCTCTACACCTGAGTCAGGCTGTCAGGCTGTCAGGAAACTTCCACAGTAT	1351
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366	sppheIlePhetYrsValLeuIvalGlyLysAlaAlaAlaIth-GlySer	382
11402	ATTCAATTCTCATAGCTGCTGCTGGTAAAGCCTCACAGCCAGT	1451
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11452	GGGACTGGAAACACAACCATAAGCTGTTGTAGCATATAATGGTTT	1501
399	uCysLeuThrLeuLeuLeuAlaIphelysLysAlaLeuProAlaL	416
11502	GGGCTTACATTATACIICCTGCGATTTCAAAGAATGGCAGCTC	1551
416	euroProIleSerIlePhetYleuIlePhetYPheserThrAspAsn	432
11552	TTCCATATCCTCACCCATTGGCTTGTCTACTTGGCACAGATTAT	1601
433	LeuValArgProPheMetAspThrLeuLaserHisGlnLeuTyryle	448
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documentation_block:		
V29525;	standard; cDNA; 2764 BP.	
13-OCT-1998	(first entry)	
Homo sapiens	PS-1 cDNA.	
PS-1; presenilin; presenilin-1; PSP-1; Alzheimer's disease; serine protease; neurodegeneration; predisposition; diagnosis		
Homo sapiens.		

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seq_documentation_block:
ID T59535; standard; cDNA; 1762 BP.
AC T59535;
DT 07-MAY-1987 (first entry)
DE Human early onset Alzheimer's disease (EOAD) gene
Early onset Alzheimer's disease; EOAD; neurodegenerative disease;
diagnosis; gene therapy; antisense; ds.
Homo sapiens.
Key OS Homo sapiens.
FT FH
variation FT FT FT FT FT FT FT FT FT PN PN
W09703086-A1.
FT PN PN
30-JAN-1987.
PP 26-JUN-1986; U11064.
PR 13-JUL-1985; US-001142.
PR 18-JUL-1985; US-001501.
(UYSF ) UNIV SOUTH FLORIDA.
Hardy JA;
PR WPI: 97-118980/11.
DR P-PSDB; W11839.
PT Early onset Alzheimer's disease gene - useful for diagnosing a
pre-disposition to Alzheimer's disease
CC Claim 1; Fig 1; 44p; English.
CC A full-length cDNA (T59535) of the early onset Alzheimer's disease
(CC) (EOAD) gene sequence codes for a 467-amino acid polypeptide (W11839).
CC Another full-length cDNA (T59536) of an EOAD splice variant gene
(CC) codes for a 463-amino acid polypeptide (W11840). The 2 sequences
(CC) can be used to generate primers and probes for the diagnosis of
(CC) predisposition to Alzheimer's disease, esp. EOAD. They can also be
(CC) used for prodn. of EOAD polypeptides in transformed host cells, and
(CC) antisense sequences can be used for the treatment of EOAD.
Sequence 1762 BP; 478 T;
SQ 442 A; 430 C;
389 C; 478 T;

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alignment_scores:
    Quality: 1454.00
    Ratio: 3.9.19
    Percent_Similarity: 79.614
    Length: 466
    Gaps: 9
    Percent_Identity: 65.451
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alignment-block:  
US-08-509-359B-137 x T59535 ..  
Alian sea 1/1 to: T59535 from: 1 to: 1762
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24 GluSerProThrProArgSerCysGlnGluGlyArgGlnGlyProGluAs 40
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 180 GAGTTTACCGTCACCGTCTACTTCCGAATCACAGATGTGAGGA 229

40 pGly GluAsnThrAlaGlnTrpArgSerGlnGluAsnGluGluA 55
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 229

74
...GLYARGPROPGLY

327 AATGGCGACCCAGGTAACTCCCCAGGTGAGCAAGTGGCA 376
 75 .LeuGluGluLeuThrLeuLysTyrGlyAlaLysValleMetL 91
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 377 AGAAAGTGAGGAGCAGACATTGAAATATGGGCCAAGCATGATCATGC 426

337 roGluValPheGluProProLeuThrGlyTyrProGly..... 349
1171 GTGAGGAATGGAAAGCCAGACGGCTCATGCCTCATGCCTCT 1220
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1221 ACACCTGAGTCACGAGCTGGTGTCAAGGAACATTCCAGCAGTATCCCGC 1270
353GluGluGluGluGluArgGlyValLysLeuGlyLeuGlyAspPheI 368
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368 lePheTyrSerValLeuValGlyLysAlaAlaAlaAlaThrGlyAsp 384
1321 TTTTCAACAGNGTTCAGGTTGGTAAGGCCCTAGCAACAGCCAGTGAGAC 1370
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1371 TGGAAACAACCATAGCCGTGTTGAGCCATATATGGTTTGCT 1420
401 uthrLeuLeuLeuAlaValPheIysLysAlaLeuProAlaLeuProI 418
1421 TACATATTACTCCATTCCATTTCAGAAAGCATGGCCAGCTCTCCAA 1470
418 leSerIleThrPheGlyLeuIlePheTyrPheSerThrAsnLeuVal 434
1471 TCTCCATCACCTTGSCCTTGSCCTTGCTTACTTGTGCACAGATTATGTGA 1520
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 416 GACGCCCTTAATGGCCGAAAGCCCCACGCCGCTCAGCAGGAGG 465
 34 lyArgGlnGlyProGluAspGlyGluAsnThrAlaGlnTrpArgSerGln 50
 466 CGAGCAGGGCCCAAGGGATGAGAGATACTGCCCACTGGAAAGCCAG 515
 51 GluAsnGluGluAspGlyGluGluAspProAspArgTyroValCysSerG1 67
 516 GAGAACGAGGAGCACGTGAGGAGGACCCCTGACCGCTATGTCGTGAGTGG 565
 67 yva!ProLyArgProProGlyLeuGluGluLeuThrLeuLysTrg 84
 566 GTTCCCCGGCGCCGGCAGGGTGGGAAAGTGACCTCAAAATACG 615
 84 LysAlaHisValIleMetLeuPheValProValThrLeuCysMetIle 100
 616 GAGCGAACCATGTGATCATCGTGTTCATGCGTCACTCTGTCATGTC 665
 101 ValAlaValAlaThrIleLysSerValArgPheTerThrGluLysAsnG1 117
 666 GTGGTGTAGGCCAACATCAAGCTGTCGGCTCTACAGAACGAAATGG 715
 117 ycinLeuIleTerThrProPheTerGluAspThrProSerValGlyIna 134
 716 AACGCTATCTACGCATTCATGAGAACACCCCTGGGGGCCAGC 765
 134 rgLeuLeuAsnSerValLeuAsnThrIleLeuMetIleSerValIleVal 150
 766 GCCCTCTCAACTCCGTCGTTGAAACCCCTCATGATGACGCTCATCGT 815
 151 ValMetThrIleLeuValLeuTerLysTerArgCystTerPheTerIle 167
 816 GTTATGACCATCTCTCTGGTGTGCTCTACAGTAGCTGGCTCATCAGT 865
 167 eileHisGlyTerPheLeuIleMetSerLeuMetLeuLeuPheLeuPheTer 184
 866 CATCAGGGCTGTGACATGCTTACTGTGCTCTGTCCTCTCA 915
 184 hrTyTrIleTerLysTerLysTerLysTerAsnValAlaMetAsp 200
 916 CCTATATCTACCTGGGAAGCTGCCTCAAGACCTACAATGTGGCATGAC 965
 201 TyrProThrLeuLeuLeuTerLysTerArgCystTerPheTerIle 217
 966 TACCCCAACCTCTGGCTGACTGTGAAACTCGGGCAGTGGCANGT 1015
 217 IcysIleHistDPrLysGlyProLeuValGlnGlnAlaTerLeuIle 234
 1016 GNGCATCACTGGCCCTCTGGCTGAGCAGGCCATCTCATCA 1065
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 1066 TGATCAGTGCGCTCATGCCCTAGTGTCAAGTAGCTCCAGAAGTGG 1115
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 1116 TCCGGGCGGGCTCATCTGGGCCATCTGTCATGATCTGGCTGT 1165
 267 IleuCysProLysGlyProLeuArgMetLeuValGluThrAlaGlnGluA 284
 1166 GCTGTGCCCCAAGGGCTCTGAAATGCTGGTAAGACTGCCAGGAGA 1215
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1216 GAATGAGGCCATATTCCCTGATATACTCATCTGCATGGTGG 1265
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 1266 ACGGTTGGATGGGAAGCTGCACCCCTCTCTCAGGGTCGCCCTCAGCT 1315
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 seq_name: /cgn2_6/ptodata/2/ina/5D_COMB.seq:US-08-875-972-28
 seq_documentation_block:
 ; Sequence 28, Application US/08875972
 ; Patent No. 5985564
 ; GENERAL INFORMATION:
 ; APPLICANT: Huntington Potter and Jimhue Li
 ; TITLE OF INVENTION: ASSAY FOR IDENTIFYING GENES CAUSING
 ; NUMBER OF SEQUENCES: CHROMOSOME NON-DISJUNCTION
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: HAMILTON, BROOK, SMITH & REYNOLDS, P.C.
 ; STREET: Two Militia Drive
 ; CITY: Lexington
 ; STATE: Massachusetts
 ; COUNTRY: USA
 ; ZIP: 02173-4799
 ; COMPUTER READABLE FORM:
 ; COMPUTER TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patient Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/875,972
 ; FILING DATE: 08-AUG-97
 ; CLASSIFICATION: 435
 ; PRIORITY APPLICATION DATA:
 ; APPLICATION NUMBER: US 60/002,448
 ; FILING DATE: 16-AUG-1995
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Granahan Esq., Patricia
 ; REGISTRATION NUMBER: 32,227
 ; REFERENCE/DOCKET NUMBER: HU95-03PA
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (781) 861-6240

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/875,972
 FILING DATE: 08-AUG-97
 CLASSIFICATION: 435
 PRIORITY APPLICATION DATA:
 APPLICATION NUMBER: US 60/002,448
 FILING DATE: 16-AUG-1995
 ATTORNEY/AGENT INFORMATION:
 NAME: Granahan Esq., Patricia
 REGISTRATION NUMBER: 32,227
 REFERENCE/DOCKET NUMBER: HU95-03PA
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (781) 861-6240
 TELEFAX: (781) 861-9540
 SEQUENCE CHARACTERISTICS:
 LENGTH: 1417 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: double
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 2..1129
 US-08-875-972-1

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 Ratio: 4.861 Gaps: 2
 Percent Similarity: 91.257 Percent Identity: 89.071
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 55 AGGGAAAGCACGCGATCATGCTGTGCGCTGACCTGATGATGATCG 104
 101 alValValAlaThrIleLysSerValArgPhenylTyrLysAsnGly 117
 105 TGTGTTAGCCACATCAAGTCAGTCGTGCCCTTCACACAGAGAAATGGA 154
 118 GluLeuIleTyrThrProPheIleGluAspThrProSerValGlyGlnAr 134
 155 CAGCTCAACTAACGCCATTCAAGTCAGTCGTGCCCTGCGTGCAGCC 204
 134 gIleLeuAsnSerValLeuAsnThrLeuIleMetIleSerValIleVal 151
 205 CCTCTCAACTCAGTCGTGCCCTCAAGTCAGTCGTGCCCTGCGTGCAGCC 254
 151 alMetThrIlePheIleValValIleTyrLysTyrLysIleTyrLysIle 167
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 168 IleHisGlyTrpIleLeuMetSerSerLeuMetIleLeuPheLeuPheIle 184
 305 ATCCATGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 354
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 seq_documentation_block:
 ; Sequence 135, Application US/08967101.
 ; GENERAL INFORMATION:
 ; APPLICANT: ST. GEORGE-HYSLOP, PETER H
 ; APPLICANT: ROMMENS, JOHANNA M
 ; APPLICANT: FRASER, PAUL E
 ; TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 ; TO ALZHEIMER'S DISEASE
 ; NUMBER OF SEQUENCES: 183
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: TESTA, HURWITZ & THEBEAULT
 ; STREET: High Street Tower - 125 High Street
 ; CITY: Boston

292	ACCGAGCATGACAGGGAGAGACTGACAACCTTGGCCAAATCTAA	341
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1092	CCGAAGGCCAA.....AGGAGGGTACCCAAGAACCCCCAAAGTAAACAC	113
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seg_documentation_block:
; Sequence 3, Application US/08670964
; Patent No. 6010874
; GENERAL INFORMATION:

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 1471 TCTCACATCACCTTGCCCTACTTGTGTTCTGAGATTATCTGTA 1520

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 1521 CAGCTTATTATGGACCAATTAGCTTCAATTATC 1562

seq_name: /cgn2_6/podata/2/ina/5D_COMB.seq:US-08-670-479-24

seq_documentation_block:

; Sequence 24, Application US/08670479

; GENERAL INFORMATION:

; APPLICANT: Goate, Alison M.

; HAROLD, John A.

; TITLE OF INVENTION: MUTANT S182 GENES

; NUMBER OF SEQUENCES: 24

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: SmithKline Beecham Corporation

; STREET: 709 Swedeland Road

; CITY: King of Prussia

; STATE: PA

; ZIP: 19406-0339

; COMPUTER READABLE FORM:

; COMPUTER: IBM Compatible

; OPERATING SYSTEM: DOS

; SOFTWARE: FASSEQ Version 1.5

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/670,479

; FILING DATE: 26-JUN-1996

; CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 60/001,500

FILING DATE: 18-JUL-1995

APPLICATION NUMBER: 60/001,800

FILING DATE: 02-AUG-1995

ATTORNEY/AGENT INFORMATION:

NAME: Han, William T.

REGISTRATION NUMBER: 34,344

REFERENCE/DOCKET NUMBER: P50361

TELECOMMUNICATION INFORMATION:

TELEPHONE: 610-270-5219

TELEFAX: 610-270-5090

TELEX:

SEQUENCE CHARACTERISTICS:

LENGTH: 1914 base Pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

HYPOTHETICAL: NO

ANTI-SENSE: NO

FRAGMENT TYPE:

ORIGINAL SOURCE:

FEATURES:

US-08-670-479-24

alignment_scores:
 Quality: 1449.00 Length: 464
 Ratio: 3.916 Gaps: 8
 Percent Similarity: 79.741 Percent Identity: 65.086

alignment_block:
 US-08-670-479-24

Align seg 1/1 to: US-08-670-479-24 from: 1 to: 1914

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 617 CCATCATGATCAGCTGCTGCTGCTGCTCATACTAACCTCTGGCTTC 666
 160 TyrLysTerGlyCystYllySpheIleHisGlyTrpLeuIleMetSerSe 176
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 967 TCAGTATATGATTAGTGGCTGTTGCTGCTGCTGCTGCTGCTGCTG 1016

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310	SerSerGlnGlyAlaLeu.....GlnLeuProtryptophanylProSerI	323
1455	GGAACTCTAAAGGAGATATCCAAGATTAATGCAAGAAGGAC	1504
323	tGluGluAspSerTyrAspSerPheGlyGlu.....ProSerI	337
1505	AGAAAGGGACTCACAGACACTGTTCAAGAGATGTGATGGGGGTTCA	1554
337	roGluValPheGluProProLeuThrGlyTyrProGly.....	349
1555	GTGAGGAATGGGAAGCCAGAGGGAGCCTCTAGGGCCCTCATCGCTCT	1604
350GluGluLeu.....GluGluLeu.....	352
1605	ACACCTGAGTCACGAGCTGTCGTCAGGAACATTCAAGCAGATCTCGC	1654
353	GluGluGluGluGluIargGlyValLysLeuGlyLeuGlyAspPheI	368
1655	TGGTGAAGACCCAGAGAAAGGGACTAAACTGGTGGGAGATTICA	1704
368	lePheTyrsValLeuValGlyLysAlaAlaLhrGlySerGlyAsp	384
1705	TTCATCTACAGTGTCTGGTGTGAGCCATAGCAAGCCAGTGAGAC	1754
385	TrpAsnThrThrLeuIaLysPheValIaLleLeuIleGlyLeuCysL	401
1755	TGGAAACACACCATAGCCTGGTCAAGAAACCATTTGAGCCATTATTGTTG	1804
401	uThrLeuLeuLeuAlaValPhenylsLysAlaLeuProAlaLeuProI	418
1805	TACATTATTAATCTCCTTGCCATTTCAGAAACCATTTGAGCTCTCAA	1854
418	leSerIlePheGlyLeuIlePheTyrsSerThrAspAsnLeuVal	434
1855	TCCUCATCACCTTGGCTGTTGTTACTTGCCACAGATTTATCCTTCA	1904
435	ArgProPheMetAspPheIleLeuAlaSerHisGlnLeuItyrile	448
1905	CAGCTTITATGGCAAAATTAGCATCATTTATCATTATATC	1946
seq_name:	/cgn2_6/ptodata/2_ina/5D_COMB.seq	US-08-592-541-5

OM of: US-08-509-359B-137 to: EST:* out_format : pfs
 Date: Mar 18, 2000 11:52 PM

About: Results were produced by the GenCore software, version 4.5.
 Copyright (c) 1993-1998 CompuGen Ltd.

Command line parameters:
 -MODEL=frame p2n.model -DEV=x1D
 -Q=>/Cgn2_1/USPTO_spool1/US08039359/runct_17032000_164054_19458/app_query.fasta.1
 -DB=EST -QFMT=fastap -SUFFIX=rst -GAPOF=12.000 -GAPEXT=4.000
 -MINMATCH=0.100 -LOOPCL=0.000 -LOOPEXT=0.000 -QAPOF=4.500
 -QGAPEXT=0.050 -XGAPOF=10.000 -XGAPEXT=0.500 -FGAPOF=6.000
 -FQGAPEXT=7.000 -YQAPOF=7.000 -DEQOF=6.000
 -DEQEXT=7.000 -START=1 -MATRIX=blosum62 -TRANS=human40 cddi
 -LIST=45 -DOCALIGN=200 -THR_SCORE=pct -ALIGN=15 -MODE=LOCAL
 -NCFMT=pfs -NORM=ext -MINLN=0 -MAXLN=1000000 -USER=US08509359
 -NCPU=6 -ICPU=3 -NO_XLXY -WAIT -THREADS=1

Search information block:
 Query: US-08-509-359B-137
 Query length: 448
 Database: EST:
 Database sequences: 4538634
 Database length: 1887831982
 Search time (sec): 350.460000

seq_name: qb_est36;AI925372	seq_documentation_block:
LOCUS	AI925372 703 bp mRNA
DEFINITION	wn53d06 x1 NCI-CGAP_Lu19 Homo sapiens cDNA clone IMAGE:2449163 3'
ACCESSION	EST similar to SW:PSN1_HUMAN P49810 PRESENILIN 2 ; mRNA sequence.
VERSION	AI925372
KEYWORDS	EST.
SOURCE	ORGANISM
	Homo sapiens
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
	Butheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE	1 (bases 1 to 703)
AUTHORS	NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE	National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL	Unpublished (1997)
COMMENT	On May 18, 1998 this sequence replaced gi:3137011. Contact: Robert Strausberg, Ph.D. Tel: (301) 496-1550
RISSUE	Email: Robert.Strausberg@nih.gov
Procurement	Emmert-Buck, M.D., Ph.D.
Preparation	M. Bento Soares, Ph.D.
CDDNA Library	CDNA Library Arrayed by: Greg Lannon, Ph.D.
DNA Sequencing	Sequencing by: Washington University Genome Sequencing Center
Clone distribution	Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LINN at: www-bio.linni.gov/bbrp/image.html
Seq_primer:	-40UP from Gibco
	High quality sequence stop: 459.
Location/Qualifiers	Source
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	/db_xref="taxon:9606"
	/clone="IMAGE:2449163"
	/clone_id="NCI-CGAP_Lu19"
	/tissue_type="squamous cell carcinoma, poorly differentiated (4 pooled tumors, including primary and metastatic)"
	/dev_stage="adult"
	/lab_host="DH10B (phage-resistant)"
	/note="Organ: lung; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from pooled lung tumor tissue, and was then primed with a Not I - Oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and Eco RI sites of the modified pT7T3 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."
FEATURES	BASE COUNT
	138 a 203 c 177 g 184 t 1 others

score_list:	Strd Orig	ZScore	EScore	Len	! Documentation
Sequence	Strd36;AI925372	55.00	234.25	8.2e-122	703
qb_est31;AI675803	+ 1120.00	2273.18	7.5e-118	661	! AI925372 wn53d06 x1 NCI-CGAP_Lu19
qb_est35;AI831581	+ 1118.00	2266.72	1.3e-117	680	! AI675803 w97a12.x1 NCI-CGAP_P1
qb_est39;AI93681	+ 910.00	208.8	3.3e-104	587	! AI831581 w39d0.x1 NCI-CGAP_Lu19
qb_est39;AW131752	+ 916.00	1855.74	1.3e-94	597	! AA993681 ot37b02.s1 Soares_tot0
qb_est17;AA62396	+ 890.00	1803.65	1.1e-91	538	! AW131752 xf34b09.x1 NCI-CGAP_B1
qb_est33;AI29395	+ 877.00	1777.55	1.0e-90	513	! AA602396 no0d05.s1 NCI-CGAP_P1
qb_est33;AI765870	- 870.00	1759.27	3.2e-89	707	! AA1829395 wf48f10.x1 NCI-CGAP_P1
qb_est44;AW12169	- 847.00	1715.02	9.2e-87	555	! AW1765870 wf66e09.x1 NCI-CGAP_P1
qb_est8;AI563205	+ 824.00	1659.65	2.0e-85	591	! AW212769 wf66e03.x1 NCI-CGAP_Ma
qb_est25;AI1276606	+ 824.00	1659.57	3.1e-84	481	! AA0563205 q132a05.s1 Soares_ret1
qb_est22;AI1046485	+ 739.00	1494.79	1.7e-74	494	! AI276606 q171h07.x1 Soares_NHh
qb_est19;AI7396	+ 732.00	1411.90	8.9e-74	476	! AI046485 ud0d05.y1 Sugano_mous
qb_est23;AI997783	+ 704.00	1423.23	1.5e-70	483	! T03796 TB913 Infrant brain. Bent
qb_est36;AI927349	+ 704.00	1445.15	1.3e-70	413	! AI097783 wf25g10.y1 Sugano_mous
qb_est24;AI25596	+ 671.00	1354.72	1.1e-66	515	! A1927349 wn9f05.x1 NCI-CGAP_Lu19
qb_est11;AA260597	+ 663.50	1339.42	7.7e-66	511	! A12225996 w108a11.y1 Sugano_mous
qb_est22;AI039465	+ 651.00	1316.98	1.4e-64	393	! AA260597 wf26g09.r1 Soares_mous
qb_est44;AW177499	+ 651.00	1314.92	1.8e-64	465	! AI039465 ox40a06.s1 Soares_tot0
qb_est44;AW177510	+ 650.00	1312.82	2.3e-64	467	! AW177499 pm4-Ct0155-770899-001-01
qb_est20;AA862334	+ 621.00	1255.76	3.5e-61	382	! AA862334 og4a05.s1 NCI-CGAP_K1
qb_est11;AA232231	+ 612.00	1228.29	3.3e-60	352	! AA782235 a131c05.s1 Soares_para
qb_est14;AA451661	+ 591.00	1193.35	1.1e-57	409	! AA451661 zx43f07.r1 Sores_tot0
qb_est16;AA537785	+ 555.00	114.97	4.4e-53	591	! AA537785 vik5e03.r1 Soares_mous
qb_est11;AA265273	+ 475.00	956.11	1.7e-44	382	! AA268820 vb01c10.r1 Soares_mous
qb_est25;AI324363	+ 539.50	1086.90	8.9e-52	435	! AI324363 ml15d23.y1 Soares_mous
qb_est11;AA237206	+ 534.00	1074.67	4.3e-51	470	! AA237206 mx18e01.r1 Soares_mous
qb_est5;N27802	+ 474.00	950.50	3.5e-44	323	! AA237206 mx18e01.r1 Soares_mous
qb_est26;AA811957	- 474.00	950.50	3.5e-44	511	! AR811957 UI-R-A09-f-06-01
qb_est10;AA144382	+ 473.50	950.53	3.5e-44	469	! AA144382 mr15d12.r1 Soares_mous
qb_est28;AI24576	+ 472.00	950.48	3.5e-44	366	! A1287167 ui73h11.y1 Sugano_mous
qb_est11;AA265273	+ 478.50	961.96	8.1e-45	426	! R1464500 y34ab1.r1 Soares_fetal
qb_est5;N27802	+ 456.00	913.88	9.9e-44	432	! A1528670 mr15d12.x1 Soares_mous
qb_est2;AA144382	+ 455.50	928.53	2.1e-42	313	! R059079 ui73h10.r1 Soares_mous
qb_est11;AA231081	+ 443.00	845.04	3.4e-38	642	! AA231081 mw1ld1.r1 Soares_mous
qb_est2;RI4600	+ 470.50	949.59	4.0e-37	306	! R1464500 y34ab1.r1 Soares_fetal
qb_est28;AI258670	- 469.50	943.32	9.9e-44	432	! A1528670 mr15d12.x1 Soares_mous
qb_est2;AA144382	+ 455.50	913.88	3.9e-42	313	! R059079 ui73h10.r1 Soares_mous
qb_est11;AA210480	+ 417.50	834.50	1.0e-37	513	! AA210480 mo86b02.r1 Beddington
qb_est18;AA673862	+ 413.00	824.06	3.9e-37	566	! AA673862 vo82h01.r1 Barstead

Align seg 1/1 to: AI925372 from: 1 to: 703
 79 LeuThrLeuLysTyrGlyAlaLysHisValLeuMetLeuPheValProVa 95

/note="Organ: lung; Vector: pTT73D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from pooled lung tumor tissue, and was then primed with a Not I primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT	137 a	191 C	173 g	178 t	1 others
ORIGIN					
alignment_scores:					
Quality: 1118.00	Length: 226				
Ratio: 5.059	Gaps: 0				
Percent Similarity: 97.788	Percent Identity: 96.903				
alignment_block:					
US-08-509-359B-137 x AI831581 ..					
Align seg 1/1 to: AI831581 from: 1 to: 680					
79 LeuthrIleuLysTyrGlyAlaLysHisValIleMetLeuPheValProva 95					
: : : : : : :					
3 CTGACCCCTCAAATACTGTGAGCCTGATCGACCTGATGCCCTGCCT 52					
95 1ThrLeuCysMetIleValValAlaLalaThrIleLyssSerValArgPheT 112					
: : : : : : :					
53 CACTCTTGTCATGATCATTGCTGGCTTAATGCCCTGCCCTGAATACTC 102					
112 YrrThrGluLysAsnGlyIleNleIleIleYrrThrProPheThrGluAspThr 128					
: : : : : : : :					
103 ACACAGGAAAGAATGGACGACTGCTACGCTACATCTGAGCACACA 152					
129 ProSerValGlyIleNleIleAsnSerValLeuAsnThrIleIle 145					
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153 CCCTCGGGGCCAGGCCATCCCTCAACTCCTGCTGTAACACCCTCATCAT 202					
145 triSerValIleValValMetThrIlePheLeuValLeuThrIleLysT 162					
: : : : : : :					
203 GATCAGGGTCATCGTGGSATGACCATTTCTGGTGSCTGCTAACAGT 252					
162 YrArgCystTyrLysSheIleHisGlyIleLeuIleMeteSerSerLeuMet 178					
: : : : : : :					
253 ACGCTGTACAAGTTCACTATGCCCTGGCTGTGATCATGCTTCACTGATG 302					
179 LeuLeuPheThrTyrIleTyrLeuGlyIleValLeuLeuLysThr 195					
: : : : : : :					
303 CGCTGTCTTCATCCATCTACCTATCTACCTTGGGAAGCTCTAACACCTA 352					
REFERENCE 1. (bases 1 to 680)					
AUTHORS NCI-CAP					
DEFINITION http://www.ncbi.nlm.nih.gov/ncicgap .					
LOCUS w139g04.x1					
VERSION P49810 PRESENILIN 2 ; , mRNA sequence.					
ACCESSION AI831581.1					
KEYWORDS EST.					
SOURCE Human.					
ORGANISM Homo sapiens					
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;					
Primates; Catarrhini; Hominidae; Homo.					
REVISION 1 On Dec 20, 1995 this sequence replaced gi:1130797.					
COMMENT Contact: Robert Strausberg, Ph.D.					
TELEPHONE (301) 496-1550					
EMAIL: Robert.Strausberg@nih.gov					
TISSUE PROCUREMENT: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.					
CODNA LIBRARY PREPARATION: M. Bento Soares, Ph.D.					
DNA SEQUENCING BY: Greg Lennon, Ph.D.					
CODNA LIBRARY ARRANGED BY: Washington University Genome Sequencing Center					
JOURNAL Tumor Gene Index					
COMPLEMENT Unpublished (1997)					
CONTACT: Robert Strausberg, Ph.D.					
LOCATION/QUALIFIERS High quality sequence stop: 456.					
1. .680 Location/Qualifiers					
/organism="Homo sapiens"					
/db_xref="taxon:606"					
/clone="IMAGE:2405191",					
/clone_lib="NCI-CGAP clone Ju19"					
/tissue_type="squamous cell carcinoma, poorly differentiated, (4 pooled tumors, including primary and metastatic)"					
/dev_stage="adult"					
/lab_host="DB10B (phage-resistant)"					
FEATURES 295					
SOURCE					
/clone="IMAGE:2405191",					
/clone_lib="NCI-CGAP clone Ju19"					
/tissue_type="squamous cell carcinoma, poorly differentiated, (4 pooled tumors, including primary and metastatic)"					
/dev_stage="adult"					
/lab_host="DB10B (phage-resistant)"					

FEATURES	High quality sequence stop: 397.
source	Location/Qualifiers 1. 597 /organism="Homo sapiens" /db_xref="Taxon:9606" /clone="IMAGE:2619929" /clone.lib="NCI_CGAP_Brn50" /tissue_type="medulloblastoma" /lab_host="DH10B (phage resistant)"
	/note="Organ: brain; Vector: pRT3D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from medulloblastoma tumor tissue, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pRT3 vector. This library is normalized. Library constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT	119 a 170 c 147 g 161 t
ORIGIN	Align seg 1/1 to: AW131752 from: 1 to: 597
alignment_scores:	Qual: 916.00 Length: 199 Ratio: 4.796 Gaps: 0 Percent Similarity: 95.980 Percent Identity: 92.462
alignment_block:	US-08-509B-137 x AW131752 ..
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	IThrLeuLysMetIleValValValAlaThrIleLeuSerValArgPhePhe 112 : 51 CACTCTGGCATATCGTGGCTAGCCACCATCAAGTCGTGCCCCCTCT 100
	YThrGluLysAsnGlyGlnLeuIleItyThrProPheThrGluAspThr 128 : 101 AGCAGAGAAGATGGAAGCTCATCTACGCCATTCTGAGGACACA 150
	ProSerValGlyInArgLeuLeuAsnSerValLeuAsnThrLeuIleMe 145 : 151 CCTCTGGGGCCAGGCCTCCATCACCCATCATCTGAAACCCCTCAT 200
	TleSerValIleValValMetThrIlePheLeuValLeuThrValIleT 162 : 201 GATCAGCTCATGGGGPATGACCATCTCTCTGGCTGCTCAAGT 250
	YArgCystyrLysPheIleHisIlyTrpLeuIleMetSerSerLeuMet 178 : 251 ACGGCTGTACAGNTCATCCATCCTGCTGATGTCATGCTTCATGAG 300
	LeuLeuLeuLeuPheThrTyryIleTrpGlyGluValLeuLysThrI 195 : 301 CTGCTGTCCTCTCACCATATCACCTTGGAAGTCCTCAAGACCTA 350
	AspValIalaMetAspThrProLeuLeuLeuThrValIlePhePheG 212 : 351 CAATGTGCCATGACTACCCCACCTCTGCGACATCTGTCGGAACTTG 400
	IyAlaValGlyMetValCysIleHistPlysGlyProLeuValIleGin 228 : 401 GGGCAGTGCGGCTGTCATGTCATCCACTGAAAGGCCCTCTGGCTGCG 450
	GinAlaArgLeuIleMetIleSerAlaLeuMetAlaLeuValPheIleE 245 : 451 CACGCCTACCTCATCATGCGTCATGCTCATGTCATGTCATCAA 500
	StryLeuProGluItpSerAlaPraValIleLeuGlyAlaIleSerVal 262 : 94 ProValThrLeuCysMetIleValValValAlaThrIleValValVal 110

DEFINITION wh66e09_x1 NCI_CGAP_Kid11 Homo sapiens cDNA clone IMAGE:2385736 3' similar to SW:PSN2_HUMAN P49810 PRESENTIN 2 ; mRNA sequence.

ACCESSION AI765870
VERSION AI765870.1
KEYWORDS EST.
SOURCE human
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 707)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT On Feb 17, 1998 this sequence version replaced gi:2889754.
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
DNA Library Preparation: M. Bento Soares, Ph.D.
DNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LINN at:
www-bio.llnl.gov/bbrp/image/Image.html

FEATURES source

Seq Primer: -40UP from Gibco
High quality sequence stop: 466.
Location/Qualifiers 1..707

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone=IMAGE:2385736"
/clone_lib="NCI_CGAP_Kid11"
/lab_host="DH10B"

/note="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA from the normalized library NCI_CGAP_Kid3 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The tracer was PCR amplified cDNAs from a pool of 5,000 clones made from the same library (clone IDs 1322376..1323911, 1456007..1456775, and 1500552..1502835). Subtraction by Bento Soares and M. Fatima Bonaldo."
Align seg 1/1 to reverse of: AI765870 from: 1 to: 707

BASE COUNT ORIGIN 177 a 197 c 182 g 150 t 1 others

alignment_scores:
Quality: 870.00 Length: 186
Ratio: 4.807 Gaps: 0
Percent Similarity: 97.312 Percent Identity: 95.161

alignment_block:
us-08-509-359B-137 x AI765870/rev ..

Align seg 1/1 to reverse of: AI765870 from: 1 to: 707

263 AsparteLeuAlaValLeuCysProLysGlyProLeuArgMetLeuValG1 279
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
707 GATCTGGGGCTGTGTGTCAAGGGTTCTGAAAGTGCTGTAGA 658
279 uThraLaGlnGluArgAsnGluProLeuPheProAlaLeuLeuTyrsers 296
657 AACTGCCAGGAAGAGAAATGAGCCCATATCCCTGCCTGTATAA_NCAT 609
296 erAlaMetLeuTrpIleValGlyMetAlaLysLeuLysProSerSerGln 312
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
608 CTGGCATGGTGTGGCTGGATGGTGGCATGGAAANGCTGACCCCTCC..TCCTCAG 560
313 GlyAlaLeuGlnLeuProTyrasProGluMetGluGluAspSerTyras 329

559 GGTGCCCCCTCACGGTCCCCTAACCGCCGAGATGGAAAGAACCTCTATA 510
329 pSerPheGlyGluProSerTyproGluValPheGluProProLeuThrG 346
509 CAGTTTGGGGCCCTCATACCCGAGTCTGCTTGAAGCTCCCTTGACTG 460
346 LysTyProGlyGluGluLeuGluGluGluGluGluGluGluGlyValLysLeu 362
459 GCTACCCAGGGAGGAGCTGGAGAAGAGAGAAAGGGCGGAGGCT 410
363 GlyLeuGlyAspPheLeuPheTyrsValLeuValGlyLysAlaAla 379
409 GGCCTCGGGACTTCATCTCATCTACAGTGTGCAGTGTGGCCAGGGCNGC 360
379 arthGlySerGlyAspThrPheAspThrLeuAlaCysHevalAlaIle 396
359 CACGGGAGGGACTGGATAACCAGCTGGCTGCTGTCATGCCATTC 310
396 euLysGlyLeuCysLeuThrLeuLeuAlaValPhelysLysAla 412
309 TCATGGCTGTTGCTGACCCCTCTGCTGCTGCTGTCAGAAAGGG 260
413 LeuProAlaLeuProLeuSerIleThrPheGlyLeuIlePheTyrsPhse 429
259 TGCCCCGCCCTCCCATTCACGTTGGGCTCATCTTTCATCTTC 210
429 rThrAspAsnLeuValArgProPheMetLaspThrLeuAlaSerHisGlnL 446
209 CACGGACACCTGGTGCCTGGCTCATTGGACACCTGCTCCATCACG 160

446 entryId 448
159 TCTACATT 152

seq_name: gb_est44:Am212769

seq_documentation_block:

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DEFINITION similar to gb:LA2177 Mus musculus S182 protein mRNA, complete cds
(MOUSE); mRNA sequence.
ACCESSION AW212769
VERSION AW212769..1 GT:6516888
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus
1..(bases 1 to 555)

REFERENCE 1
AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP).
TITLE Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT On Jul 7, 1999 this sequence version replaced gi:5406916.
Other_ESPs: u066e03.y1
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (IIML)
DNA Sequencing by: NCI-CGAP
Clone distribution: Washington University Genome Sequencing Center
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/Image.html

MGI:1027968
Seq primer: -40UP from Gibco
High quality sequence stop: 421.
Location/Qualifiers 1..555
/organism="Mus musculus"

FEATURES source

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 110 gPheTerThrGluLysAsnGlyGlnLeuIleTerThrProPheTerGluA 127
 53 CTTCTCACAGAGAAATGCACTACAGGCCATTCACTGAGG 102
 127 sPthrProSerValGlyGlnArgLeuAsnSerValLeuAsnThrLeu 143
 103 ACACACCCCTGGTGGCCAGGCCCTCACTCGTGCTAACACACCTC 152
 144 IleMetIleSerValLeuValMetThrIlePheLeuValLeuIle 160
 153 ATCATGATCACGGTCATCGTGTATGACCATCTCTGGGGTGCCTA 202
 160 rLysTerArgCystTerLysPheIleHisGlyTrpLeuIleMetSerVal 177
 203 CAAGTACCGCGCTGCTACAAGTTCATCCATGGCTGGTGTATCATGTCCTCAC 252
 177 eumetIleLeuPheLeuPheThrIleTerLeuGluValLeuIlys 193
 253 TGATGCTGCTGCTCTCTCTACCTATATCACCTGGGAAGTGTCAAG 302
 194 ThryTerAsnValAlaMetAspTerProThrIleLeuThrValTrpAs 210
 303 ACCTACATGTGGCATGGACTAACCCACCTCTCTGACTGTCTGGAA 352
 210 nPheGlyAlaValGlyMetValCysIleHistPheGlyProLeuValL 227
 353 CTTGGGGCAGTGGCATGGCTGATGGTGTGATCCACTGGAGGGCCCTCTGGTC 402
 227 euGlnGlnAlaTerLeuIleTerLeuAlaLeuAlaLeuAlaLeuAla 243
 403 TGCAGGAGGCCCTACCTCTCATGATGCTAGGGCTAGGCCCTAGGTC 452
 244 IleLeuTerLeuProGluTrpSerAlaTerPheIleLeuGlyAlaLeuSe 260
 453 ATCAAGTACTC.CCAGAGTGGTCGGTGGCTCATC...TGGGCCANC 498
 260 rValTerAspLeuValAlaValLeuCysProLysGlyProLeuArgMetL 277
 499 TCTGTGTATGATCCTGCTNGCTGTTCAAAAGGCT...CTGAGATGT 545
 277 euvalGluThrAlaGlnGluArgAsnGluPro 287
 546 GG...TGAANTGCCAGGAGAAATGAGCCA 574

seq_name: qb_est25:AI276606

seq_documentation.block:

LOCUS AI276606 481 bp mRNA EST 29-JAN-1999
 DEFINITION q11h07.x1 Soares_NHMPU_S1 Homo sapiens cDNA clone IMAGE:1877821
 ACCESSION AI276606
 VERSION AI276606.1 GI:3898808
 SOURCE
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 481)
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 tumor Gene Index
 Unpublished (1997)
 COMMENT On Jan 19, 1998 this sequence replaced gi:2285413.
 Contact: Robert Strausberg, Ph.D.
 Tel: (301) 496-1550
 Email: Robert_Strausberg@nih.gov
 This clone is available royalty-free through LiNL ; contact the
 IMAGE Consortium (infoimage.lnl.gov) for further information.
 Insert Length: 1769 Std Error: 0.00
 seq primer: -40up from Gibco

FT Domain /label= "TM3-4"
 FT Domain /note= "hydrophilic loop"
 201 . .218
 FT Domain /label= "TM4"
 /note= "transmembrane domain in 4"
 219 . .224
 FT Domain /label= "TM4-5"
 /note= "hydrophilic loop"
 225 . .244
 FT Domain /label= "TM5"
 /note= "transmembrane domain in 5"
 245 . .249
 FT Domain /label= "TM5-6"
 /note= "hydrophilic loop"
 250 . .268
 FT Domain /label= "TM6"
 /note= "transmembrane domain 6"
 269 . .387
 FT Domain /label= "TM6-7"
 /note= "hydrophilic loop"
 388 . .409
 FT Domain /label= "TM8"
 /note= "transmembrane domain 8"
 FT Misc_difference 141
 /note= "Ash141Ile mutation site (Claim 19)"
 FT Misc_difference 239
 /note= "Met239Val mutation site (Claim 19)"
 FT Misc_difference 420
 /note= "Ile420Thr mutation site"
 PN WO9801549-A2.
 PD 15-JAN-1998.
 PP 04-JUL-1997; CA0475.
 PR 02-JAN-1997; US-034590.
 PR 05-JUL-1996; US-021673.
 PR 12-JUL-1996; US-01700.
 PR 08-NOV-1996; US-029895.
 PA (HSCR-) HSC RES & DEV LP.
 (OTOR) UNIV TORONTO GOVERNING COUNCIL.
 PI Fraser PE, Rommens JM, St George-Hyslop PH;
 WPI; 98-286355/25
 DR N-PSDB; V04659.
 PT New isolated mutant presenilin-1 genes - useful for developing products for use in detection, diagnosis and therapy of Alzheimer's disease and for drug screening
 Claim 19: Page 203-204; 231pp; English.
 PS This polypeptide comprises human presenilin-2 (HPS2). Its amino acid sequence was deduced from an isolated cDNA clone (see V04659). Human and murine presenilin-1 sequences are also provided (see W23964-066). Mutations in the PS-1 and PS-2 genes are linked to the development in humans of forms of familial Alzheimer's disease (FAD) and may be causative of other disorders, e.g., cognitive, intellectual, neurological or physiological disorders such as cerebral haemorrhage, schizophrenia, depression, mental retardation and epilepsy. Use of the nucleic acids and proteins comprising or derived from the Presenilins is made in screening and diagnosing FAD, identifying and developing therapeutics for treatment of FAD, and in producing cell lines and transgenic animals useful as models of FAD. Methods for identifying substances that bind to, or modulate the activity of a presenilin protein, and methods for identifying substances that affect the interaction of a presenilin-interacting protein with a presenilin protein are also disclosed. Sequence 448 AA;
 SQ 100.0%; Score 2336; DB 1; Length 448;
 Best Local Similarity 100.0%; Pred. No. 7.3e-237;
 Matches 448; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 DRYVCSGVPGRRPGLLEELSTLKYGAKHYTMLFYPVTLCMIVVYATIKSYVRFYTEKNGQLI 120
 Db 61 DRYVCSGVPGRRPGLLEELSTLKYGAKHYTMLFYPVTLCMIVVYATIKSYVRFYTEKNGQLI 120
 FT Domain /label= "TM4-5"
 /note= "transmembrane domain in 5"
 219 . .224
 FT Domain /label= "TM4-5"
 /note= "hydrophilic loop"
 225 . .244
 FT Domain /label= "TM5"
 /note= "transmembrane domain in 5"
 245 . .249
 FT Domain /label= "TM5-6"
 /note= "hydrophilic loop"
 250 . .268
 FT Domain /label= "TM6"
 /note= "transmembrane domain 6"
 269 . .387
 FT Domain /label= "TM6-7"
 /note= "hydrophilic loop"
 388 . .409
 FT Domain /label= "TM8"
 /note= "transmembrane domain 8"
 FT Misc_difference 141
 /note= "Ash141Ile mutation site (Claim 19)"
 FT Misc_difference 239
 /note= "Met239Val mutation site (Claim 19)"
 FT Misc_difference 420
 /note= "Ile420Thr mutation site"
 PN WO9801549-A2.
 PD 15-JAN-1998.
 PP 04-JUL-1997; CA0475.
 PR 02-JAN-1997; US-034590.
 PR 05-JUL-1996; US-021673.
 PR 12-JUL-1996; US-01700.
 PR 08-NOV-1996; US-029895.
 PA (HSCR-) HSC RES & DEV LP.
 (OTOR) UNIV TORONTO GOVERNING COUNCIL.
 PI Fraser PE, Rommens JM, St George-Hyslop PH;
 WPI; 98-286355/25
 DR N-PSDB; V04659.
 PT New isolated mutant presenilin-1 genes - useful for developing products for use in detection, diagnosis and therapy of Alzheimer's disease and for drug screening
 Claim 19: Page 203-204; 231pp; English.
 PS This polypeptide comprises human presenilin-2 (HPS2). Its amino acid sequence was deduced from an isolated cDNA clone (see V04659). Human and murine presenilin-1 sequences are also provided (see W23964-066). Mutations in the PS-1 and PS-2 genes are linked to the development in humans of forms of familial Alzheimer's disease (FAD) and may be causative of other disorders, e.g., cognitive, intellectual, neurological or physiological disorders such as cerebral haemorrhage, schizophrenia, depression, mental retardation and epilepsy. Use of the nucleic acids and proteins comprising or derived from the Presenilins is made in screening and diagnosing FAD, identifying and developing therapeutics for treatment of FAD, and in producing cell lines and transgenic animals useful as models of FAD. Methods for identifying substances that bind to, or modulate the activity of a presenilin protein, and methods for identifying substances that affect the interaction of a presenilin-interacting protein with a presenilin protein are also disclosed. Sequence 448 AA;
 SQ 100.0%; Score 2336; DB 1; Length 448;
 Best Local Similarity 100.0%; Pred. No. 7.3e-237;
 Matches 448; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Query Match 1 MLTFMASDSEEVCDERTSLMSAESPRSCQEGRPQEDGENTAQWRSQENEDGEEDP 60
 Db 1 MLTFMASDSEEVCDERTSLMSAESPRSCQEGRPQEDGENTAQWRSQENEDGEEDP 60

regions. Mutations in PS genes are implicated in familial Alzheimer's disease (AD) and other diseases such as cerebral haemorrhage, schizophrenia, depression etc. so detection of mutations in the DNA encoding the wild type sequences can be used for diagnosis of these diseases. The wild type proteins, or vectors that express them or containing antisense sequences, antibodies selective for these mutant forms of the proteins and modulators of PS gene expression are potentially useful for treatment of AD etc. Transgenic animals are useful as models for drug screening. The antibodies can also be used e.g. for affinity purification and in immunoassays.

Sequence 448 AA;

Query Match	99.6%	Score 2327;	DB 1;	Length 448;
Best Local Similarity	99.8%	Pred. No.	6.4e-236;	
Matches 447; Conservative 0; Mismatches 1; Indels 0; Gaps 0;				
Db	1	MLTFMASDSEEEYCDERTSLMSAESPTPRSCQSGRQPEDGENTAQWRSQENEDGEEDP 60		
Qy	61	DRYVCSGYGPGRGPGLLEELTLKGAKHVTMFLFVTLCKMIVVATKSVRFTEKNGOLI 120		
Db	61	DRYVCSGYGPGRGPGLLEELTLKGAKHVTMFLFVTLCKMIVVATKSVRFTEKNGOLI 120		
Qy	121	YTPFIEDTPSVGQRLLNSVLNTLIMISVTVMTIFLVLYKRYCYKFHGWLIMSSLML 180		
Db	121	YTPFIEDTPSVGQRLLNSVLNTLIMISVTVMTIFLVLYKRYCYKFHGWLIMSSLML 180		
Qy	181	FIFTYIYLGEVKTYNVAMDYPFLTLLTWNFGAVGMYCITHWKGPVLQVAYLIMISALMA 240		
Db	181	FIFTYIYLGEVKTYNVAMDYPFLTLLTWNFGAVGMYCITHWKGPVLQVAYLIMISALMA 240		
Qy	241	LVEFKYLPERSAWVILGAISVYDLYAVLCPKGPRLMLVETAQERNEPFPALIYSSAMW 300		
Db	241	LVEFKYLPERSAWVILGAISVYDLYAVLCPKGPRLMLVETAQERNEPFPALIYSSAMW 300		
Qy	301	TGYNAKLDPSSQGALQPYDPNEEDSDSFGEPSYPEVFPEPLTGYPEELEEEERGV 360		
Db	301	TGYNAKLDPSSQGALQPYDPNEEDSDSFGEPSYPEVFPEPLTGYPEELEEEERGV 360		
Qy	361	KLGIGDFIFYSVLYGKAATGSGDWNTTLACFVAILIGCLTLLIAVFKALPALPISI 420		
Db	361	KLGIGDFIFYSVLYGKAATGSGDWNTTLACFVAILIGCLTLLIAVFKALPALPISI 420		
Qy	421	TFGLIIFYFSTDNLVRPFMDTLASHOLYI 448		
Db	421	TFGLIIFYFSTDNLVRPFMDTLASHOLYI 448		
RESULT	7			
W28508				
ID	W28508	standard; Protein: 447 AA.		
AC				
DT	07-DEC-1997	(first entry)		
Full AD/AD3LP sequence.				
KW				
AD3; AD4; AD3LP; Alzheimer's disease; chromosome; missegregation; presenilin; inhibitor; AD; trisomy 21; ss.				
KW				
OS				
Homo sapiens.				
PN	W0910723-A2.			
PD	27-FEB-1997.			
PR	15-AUG-1996; US-002448.			
PA	(HARD) HARVARD COLLEGE.			
PI	Li J, Potter H.			
DR	WPI: 97-165297/15.			
DR	N-PDBB; T87426.			
PT	Identifying genes which cause chromosome missegregation - useful for identifying causes of and treatments for diseases, e.g. Alzheimer's disease, cancer and ageing.			
PT	Claim 29; Fig 29; 77pp; English.			
CC	Identifying genes which cause improper chromosome segregation, screening for inhibitors of chromosome missegregation and processes			

caused by genes encoding chromosome missegregation promoters caused by genes encoding chromosome missegregation promoters given in T87401 to T87426 can be used in the above methods. It is not clear from the figure legend, the figure and the disclosure of the specific sequence which sequence of Fig 1 and Fig 28 is the AD4/AD3LP or the AD3 sequence.

Sequence 447 AA;

Query Match	99.3%	Score 2320.5;	DB 1;	Length 447;
Best Local Similarity	99.8%	Pred. No.	3.1e-235;	
Matches 447; Conservative 0; Mismatches 0; Indels 1; Gaps 1;				
Db	1	MLTFMASDSEEEYCDERTSLMSAESPTPRSCQSGRQPEDGENTAQWRSQENEEDGEEDP 60		
Qy	1	MLTFMASDSEEEYCDERTSLMSAESPTPRSCQSGRQPEDGENTAQWRSQENEEDGEEDP 60		
Db	1	MLTFMASDSEEEYCDERTSLMSAESPTPRSCQSGRQPEDGENTAQWRSQENEEDGEEDP 60		
Qy	61	DRYVCSGYGPGRGPGLLEELTLKGAKHVTMFLFVTLCKMIVVATKSVRFTEKNGOLI 120		
Db	61	DRYVCSGYGPGRGPGLLEELTLKGAKHVTMFLFVTLCKMIVVATKSVRFTEKNGOLI 120		
Qy	121	YTPFIEDTPSVGQRLLNSVLNTLIMISVTVMTIFLVLYKRYCYKFHGWLIMSSLML 180		
Db	121	YTPFIEDTPSVGQRLLNSVLNTLIMISVTVMTIFLVLYKRYCYKFHGWLIMSSLML 180		
Qy	181	FIFTYIYLGEVKTYNVAMDYPFLTLLTWNFGAVGMYCITHWKGPVLQVAYLIMISALMA 240		
Db	181	FIFTYIYLGEVKTYNVAMDYPFLTLLTWNFGAVGMYCITHWKGPVLQVAYLIMISALMA 240		
Qy	241	LVEFKYLPERSAWVILGAISVYDLYAVLCPKGPRLMLVETAQERNEPFPALIYSSAMW 300		
Db	241	LVEFKYLPERSAWVILGAISVYDLYAVLCPKGPRLMLVETAQERNEPFPALIYSSAMW 300		
Qy	301	TGYNAKLDPSSQGALQPYDPNEEDSDSFGEPSYPEVFPEPLTGYPEELEEEERGV 360		
Db	301	TGYNAKLDPSSQGALQPYDPNEEDSDSFGEPSYPEVFPEPLTGYPEELEEEERGV 360		
Qy	361	KLGIGDFIFYSVLYGKAATGSGDWNTTLACFVAILIGCLTLLIAVFKALPALPISI 420		
Db	361	KLGIGDFIFYSVLYGKAATGSGDWNTTLACFVAILIGCLTLLIAVFKALPALPISI 420		
Qy	421	TFGLIIFYFSTDNLVRPFMDTLASHOLYI 448		
Db	421	TFGLIIFYFSTDNLVRPFMDTLASHOLYI 448		
RESULT	8			
ID	W05766	standard; Protein: 414 AA.		
AC	W05766;			
DT	25-JUL-1997	(first entry)		
DE	Presenilin-2 delta26-296 mutation.			
KW	Presenilin-2; human; HPS1-1; HPS1-2; PS-2; integral membrane protein; AD; familial Alzheimer's disease; cerebral haemorrhage; schizophrenia; depression; antibody; gene expression modulator; therapy; mutein.			
KW	Homo sapiens.			
OS				
PH	Location/Qualifiers			
FT	misc_difference 263 . 264			
FT	/note= "site of 34 residue deletion"			
FT	W09634099-A2.			
PD	31-OCT-1996..			
PF	29-APR-1996.. CA0263.			
PR	28-APR-1995; US-431048.			
PR	28-JUN-1995; US-496841.			
PR	31-JUL-1995; US-509359.			
PA	(HSCR) HSC RES DEY LP.			
PA	(UTOR) UNIV TORONTO GOVERNING COUNCIL.			
PI	Fraser PE, Rommens JM, St George-Hyslop PH;			
DR	WPI: 96-4763149.			
PT	New Presenilin genes - useful for diagnosis, therapy and drug screening of familial Alzheimer's disease, cerebral disorders, etc.			
PT	Claim 4; Page -; 178pp; English.			
PS	W05763-W05766 represent mutated versions of the human presenilin-2			

coding sequence for wild type human PS-2; The presenilins are a family of highly conserved integral membrane proteins with a common structural motif, common alternative splicing patterns, and common mutational hot spot regions. Mutations in PS genes are implicated in familial Alzheimer's disease (AD) and possibly other diseases such as cerebral haemorrhage, schizophrenia, depression etc., so detection of mutations in the DNA encoding these sequences can be used for diagnosis of these diseases. These proteins, or vectors that express them or containing antisense sequences, antibodies selective for mutant forms of these proteins (such as w05736) and modulators of PS gene expression are potentially useful for treatment of AD etc. Transgenic animals are useful as models for drug screening. The antibodies can also be used e.g. for affinity purification and in immunoassays.

Sequence 467 AA;

Query Match Score 1468; DB 1; Length 467;

Best Local Similarity 64.0%; Pred. No. 9.9e-146; Mismatches 46; Indels 44; Gaps 6;

Matches 299; Conservative 46; Gaps 6;

CC coding sequence for wild type human PS-2; The presenilins are a family of

CC highly conserved integral membrane proteins with a common structural

CC motif, common alternative splicing patterns, and common mutational hot spot

CC regions. Mutations in PS genes are implicated in familial Alzheimer's

CC disease (AD) and possibly other diseases such as cerebral haemorrhage,

CC schizophrenia, depression etc., so detection of mutations in the DNA

CC encoding these sequences can be used for diagnosis of these diseases.

CC These proteins, or vectors that express them or containing antisense

CC sequences, antibodies selective for mutant forms of these proteins (such

CC as w05736) and modulators of PS gene expression are potentially useful

CC for treatment of AD etc. Transgenic animals are useful as models for drug

CC screening. The antibodies can also be used e.g. for affinity purification

CC and in immunoassays.

SQ Sequence 467 AA;

CC	FT	Domain
CC	FT	/label= TM2-3
CC	FT	/note= "hydrophilic loop"
CC	FT	155..163
CC	FT	/label= TM2-3
CC	FT	/note= "hydrophilic loop"
CC	FT	164..183
CC	FT	/label= TM3
CC	FT	/note= "transmembrane domain 3"
CC	FT	184..194
CC	FT	/label= TM3-4
CC	FT	/note= "hydrophilic loop"
CC	FT	195..212
CC	FT	/label= TM4
CC	FT	/note= "transmembrane domain 4"
CC	FT	213..220
CC	FT	/label= TM4-5
CC	FT	/note= "hydrophilic loop"
CC	FT	221..238
CC	FT	/label= TM5
CC	FT	/note= "transmembrane domain 5"
CC	FT	239..243
CC	FT	/label= TM5-6
CC	FT	/note= "hydrophilic loop"
CC	FT	244..262
CC	FT	/label= TM6
CC	FT	/note= "transmembrane domain 6"
CC	FT	263..407
CC	FT	/label= TM6-7
CC	FT	/note= "hydrophilic loop"
CC	FT	408..428
CC	FT	/label= TM8
CC	FT	/note= "transmembrane domain 8"
CC	FT	177
CC	FT	/note= "phe177Ser mutation site (Claim 1)"
CC	FT	Misc_difference 439
CC	FT	/note= "Ile439Val mutation site (Claim 1)"
CC	FT	W0801549-A.
Db	FT	PN
Db	FT	W0801549-A.
Db	FT	PD 15-JAN-1998;
Db	FT	PF 04-JUL-1997;
Db	FT	PR 02-JAN-1997; US-034590;
Db	FT	PR 05-JUL-1996; US-021672;
Db	FT	PR 12-JUL-1996; US-021700;
Db	FT	PR 08-NOV-1996; US-029895;
Db	FT	PA (HSCR-) HSC RES & DEV LP.
Db	FT	PA (UTOR) UNIV TORONTO GOVERNING COUNCIL.
Db	FT	PI Fraser PE, Rommens JM, St. George-Hyslop PH;
Db	FT	DR WPI, 98-28635/25.
Db	FT	DR N-PSDB; V0468.
Db	FT	PT New isolated mutant presenilin-1 genes - useful for developing
Db	FT	PT products for use in detection, diagnosis and therapy of Alzheimer's
Db	FT	PT disease and for drug screening
Db	FT	PT Disclosure: Page 199-00; 238pp; English.
Db	FT	CC This polypeptide comprises the murine presenilin-1 (PS1) homologue.
Db	FT	CC Its amino acid sequence was deduced from an isolated cDNA clone
Db	FT	CC (see V04668). Mutations in the human PS1 and PS2 genes (see
Db	FT	CC V0466-68) have been linked to the development in humans of forms
Db	FT	CC of familial Alzheimer's disease (FAD). All amino acids that are
Db	FT	CC mutated in analysed FAD pedigrees (see W3964) were conserved in
Db	FT	CC the murine homologue. Use of the nucleic acids and proteins
Db	FT	CC comprising or derived from presenilins can be made in screening and
Db	FT	CC diagnosing FAD, identifying and developing therapeutics for
Db	FT	CC treatment of FAD, and in producing cell lines and transgenic
Db	FT	CC animals useful as models of FAD. Methods for identifying
Db	FT	CC substances that bind to, or modulate the activity of a presenilin
Db	FT	CC protein, and methods for identifying substances that affect the
Db	FT	CC interaction of a presenilin-interacting protein with a presenilin
Db	FT	CC protein are also disclosed.
Db	FT	Sequence 467 AA;
RESULT 11	FT	Query Match 62.8%; Score 1468; DB 1; Length 467;
W23966	FT	Best Local Similarity 64.0%; Pred. No. 9.9e-146;
ID	FT	Matches 299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;
AC	FT	FT Domain
W23966	FT	W23966 standard; Protein: 467 AA.
AC	FT	AC W23966; (first entry)
W23966	FT	DE Mouse presenilin-1 homologue.
ID	FT	KW Presenilin-1; PS1 gene; mouse; familial Alzheimer's disease; FAD;
DE	FT	KW cerebral haemorrhage; schizophrenia; depression; epilepsy;
DE	FT	KW mental retardation; diagnosis; therapy; transgenic animal.
OS	FT	Mus musculus.
FH	FT	Key Location_Qualifiers
FT	FT	FT Domain 82..100
FT	FT	/note= "transmembrane domain 1"
FT	FT	101..132
FT	FT	/label= TM1-2
FT	FT	/note= "hydrophilic loop"
FT	FT	133..154
FT	FT	/label= TM2
FT	FT	/note= "transmembrane domain 2"
QY	FT	24 ESPTPRSCQEGRGPEDGENTAQWSRQNEEDGEEDPDRYVCSGP----GRPG-----75

Page 9

peptides. Antisense nucleic acid may also be used to regulate expression of the PS1/429 gene, and both nucleic acids and peptides are useful as size markers in electrophoresis, chromatography etc. The transgenic animals are used as models for AD, e.g. for testing drugs. Regulators of the PS1/429 gene or polypeptide can be used to treat e.g. AD or diseases involving mitochondrial pathology, apoptosis and neurodegeneration. Typical regulators are antisense sequences, ribozymes, aptamers, synthetic or natural compounds. (II) may also be used to target other coding sequences to particular cellular locations.

FT	peptides. Antisense nucleic acid may also be used to regulate expression of the PS1/42 gene, and both nucleic acids and peptides are useful as size markers in electrophoresis, chromatography etc. The transgenic animals are used as models for AD, e.g. for testing drugs. Regulators of the PS1/42 gene or polypeptide can be used to treat e.g. AD or diseases involving mitochondrial pathology, apoptosis and neurodegeneration. Typical regulators are antisense sequences, ribozymes, aptamers, synthetic or natural compounds. (II) may also be used to target other coding sequences to particular cellular locations.	Q	Sequence - 467 AA;
FT	Query Match 62.8%; Score 1467; DB 1; Length 467;		
FT	Best Local Similarity 65.7%; Pred. No. 1..3e-145;		
FT	Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;		
FT	Y 24 ELPAPLSYQAAQMSDNDHLSNTVRQNDNRREHQHRR-SLGHPEPLSNSRPGQNSRQ 61	b	
FT	Y 75 -----LEEEELTYKGAKHVIMLFYPTVLCKMIVVYATIKSYSTFETYEKNGQLIYTPTED 127	b	
FT	Y 62 VVEQQDEEDEELTYKGAHVIMLFYPTVLCKMIVVYATIKSYSTFYTDRGQLIYTPTED 121	b	
FT	Y 128 TPSVGORLNLNSVNLNTIMISVIVWMTLFLVLYKRYKXKFTHGLWLMISMLFLFYYIY 187	b	
FT	Y 122 TETVGORALHNTSLNAIMISVIVWMTLFLVLYKRYKXKVIAHWLISLILFFSEVY 181	b	
FT	Y 188 LGEVLRKTYNAMYDPTTLLKPLVYAVLCPKGPLRMLVYETAQERNEPIPPALIYS 247	b	
FT	Y 182 LGEVFKTYNADVYTVAWLWNVFGVGMISHWKGLRQPAYLIMISALMALVFIFYL 241	b	
FT	Y 248 PEWSAWVILGAISVSYDVLAVLCPKGPLRMLVYETAQERNEPIPPALIYSAMWTVGMAKL 307	b	
FT	Y 242 PEWTAWLILAVISVYDVLAVLCPKGPLRMLVYETAQERNETLFPALIYSTMTWLVNAEG 301	b	
FT	Y 308 DPSSQCAL---QLPYDPDE-MEEDSYDSFGE-- -PSYPFEFPEPLTGYPG----- 350	b	
FT	Y 302 DPEAQRVRSKSKYNAESTERESQDITYAENDGGFSEWEORDSHGPHRSTPESRAV 361	b	
FT	Y 350 EEL-----EEEELVYKLGDFIFIYSVLYGKAATGSDWNNTLACFCVAILIGLCT 402	b	
FT	Y 362 QELSSSILAGDPEEGVKLGDFIFIYSVLYGKAATGSDWNNTLACFCVAILIGLCT 421	b	
FT	Y 403 LLLLAVFKKALPALPISITISFGFLIFYFSTDONLYRPENDTLASHOLYI 448	b	
FT	Y 422 LLLLAVFKKALPALPISITISFGFLIFYFATDYLQPEFDQLAFTOFYI 467	b	
FT	RESULT 14		
D	W23964 standard; Protein: 467 AA.	C	
C	W23964;	C	
T	20-TTU-1998 (first entry)	T	
E	Human Presenilin-1.	E	
W	Presenilin-1; PS1 gene; human; familial Alzheimer's disease; FAD; cerebral haemorrhage; schizophrenia; depression; epilepsy; mental retardation; diagnosis; therapy; transgenic animal.	W	
S	Homo sapiens.	S	
T	Key	T	
T	Location/Qualifiers	T	
T	82..100	T	
T	/label= TM1	T	
T	/note= "transmembrane domain 1"	T	
T	101..132	T	
T	/label= TM1-2	T	
T	/note= "hydrophilic loop"	T	
T	133..154	T	
T	/label= TM2	T	
T	/note= "transmembrane domain 2"	T	
T	155..163	T	
T	/label= TM2-3	T	
T	/note= "hydrophilic loop"	T	
T	164..183	T	
T	/label= TM3	T	
T	/note= "transmembrane domain 3"	T	
T	184..194	T	
T	/label= TM3-4	T	
T	/note= "hydrophilic loop"	T	
T	195..212	T	
T	/label= TM4	T	
T	/note= "transmembrane domain 4"	T	
T	213..220	T	
T	/label= TM4-5	T	
T	/note= "hydrophilic loop"	T	
T	221..238	T	
T	/label= TM5	T	
T	/note= "transmembrane domain 5"	T	
T	239..243	T	
T	/label= TM5-6	T	
T	/note= "hydrophilic loop"	T	
T	244..262	T	
T	/label= TM6	T	
T	/note= "transmembrane domain 6"	T	
T	263..407	T	
T	/label= TM6-7	T	
T	/note= "hydrophilic loop"	T	
T	408..428	T	
T	/label= TM8	T	
T	/note= "transmembrane domain 8"	T	
T	Misc_difference 177		
T	/note= "Phe117Ser mutation site		
T	Misc_difference 439		
T	/note= "Ile439Val1 mutation site		
T	Misc_difference 257		
T	/note= "Asp257Ala mutation site		
T	residue 258-260 deletion		
T	Misc_difference 258..290		
T	/note= "residue 258-290 deletion with Asp257Ala mutation		
T	Misc_difference 143		
T	/note= "Ile143Thr mutation site		
T	Misc_difference 146		
T	/note= "Met146Leu mutation site		
T	Misc_difference 171		
T	/note= "Leu171Pro mutation site		
T	Misc_difference 260		
T	/note= "Ala260Val mutation site		
T	Misc_difference 263		
T	/note= "Cys263Arg mutation site		
T	Misc_difference 264		
T	/note= "Pro264Leu mutation site		
T	Misc_difference 267		
T	/note= "Pro267Ser mutation site		
T	Misc_difference 280		
T	/note= "Glu280Ala mutation site		
T	Misc_difference 280		
T	/note= "Glu280Gly mutation site		
T	Misc_difference 285		
T	/note= "Ala285Val mutation site		
T	Misc_difference 286		
T	/note= "Leu286Val mutation site		
T	Misc_difference 322		
T	/note= "Leu322Val mutation site		
T	Misc_difference 392		
T	/note= "Leu392Val mutation site		
T	Misc_difference 410		
T	/note= "Cys410Tyr mutation site		
T	Misc_difference 79		
T	/note= "Ala79Xaa mutation site"		
T	Misc_difference 82		
T	/note= "Val82Leu mutation site"		
T	Misc_difference 96		
T	/note= "Val96Phe mutation site"		
T	Misc_difference 115		
T	/note= "Tyr115His mutation site"		
T	Misc_difference 139		

	Matches	305;	Conservative	40;	Mismatches	79;	Indels	42;	Gaps	8;
Qy	24	ESPTPRSCQEGRQGPEDGENTAAQWRSQENEEDGEEDPDRYVGSGVP	-----	GRPPG	---	75				
Db	3	ELDAPLSYFQNQMSDNHLSITVRSONDNRREPQEHDRR-SLIGHPEPLSNGRPGNSRQ	61							
Qy	75	-----	LEEEETLKYGAKHYTMFLFYPTILCMIVVVVATIKSYRFYETKNGQLITYPTED	127						
Db	62	VYEDEEEDEETLKYGAKHYTMFLFYPTILCMIVVVVATIKSYRFYETKNGQLITYPTED	121							
Qy	128	TPEVGQRLNSVNTLIMISIVVMTTFLVVLKYRCYKFINGWLINSSMLFLFVYI	187							
Db	122	TEVGQRALHSITNAAMISIVVMTTFLVVLKYRCYKVIAWLISSLLLFFSFIV	181							
Qy	188	LGFLVLTYYNVAANDYPTULLTWNFGAYGMYC1HWKGPLVLYOOAYLIMSLALMVFTKYL	247							
Db	182	LGVEKTYNVAADYITVALLIWNFGVGMISHWKGPRLQQYLINTSLALMVFTKYL	241							
Qy	248	PENSAWVILGASVYDLYAVICPKGPRLMVETAQERNEPISPALIYSSAMMTVGMAKL	307							
Db	242	PENTAWLJLAVISVYDLYAVICPKGPRLMVETAQERNETLPFAYISSTMWLVNNAEQ	301							
Qy	308	DPSGQAL-OLPYDPE-MEEDSYDSGE--PSYPEFEPPPTGYPS-----	350							
Db	302	DPEAQRRVSKNSKYNASTERSQDTAENDGGFSEEWERQDSHIGPHRSTPESRAAV	361							
Qy	350	EEL-----EPEEEGVKLGLGDFIFYSVLYGKAATGSGDWNTLACEVALIGLCT	402							
Db	362	QEUSSIILAGEDEPEEREVKLGLGDFIFYSVLYGKASATASGDWNTTIACFVALIGLCT	421							
Qy	403	LLIAVFKKALDAPLISITGLIFYFSTDNLVPEMDTLASHOLYI	448							
Db	422	LLIAVFKKALDAPLISITGLIFYFSTDNLVPEMDTLASHOLYI	467							

Search completed: March 20, 2000, 05:31:20
 Job time: 4209 sec

Gencore version 4.5
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OM protein - protein search, using sw mode

Run on: March 18, 2000, 19:01:42 ; Search time 26.47 Seconds
 (without alignments)
 225.059 Million cell updates/sec

Title: US-08-509-359B-137
 Perfect score:
 2336
 Sequence: 1 MLTFMASDSBEVCDERTSL.....STDNLVRPFMDTLASHQIYT 448

Scoring table: BLOSUM62

Searched: 133990 seqs, 13297546 residues

Database : Issued_Patents_AA:*

Word size : 0

Number of hits that pass the threshold : 133990

1: /cgn2_6/ptodata/2/iaa/5a_COMBO.pep:*

2: /cgn2_6/ptodata/2/iaa/5b_COMBO.pep:*

3: /cgn2_6/ptodata/2/iaa/5c_COMBO.pep:*

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5: /cgn2_6/ptodata/2/iaa/backfiles1.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

ALIGNMENTS

RESULT 1
 US-08-967-101-137
 ; Sequence 137, Application US/08967101
 ; Patent No. 5840540

GENERAL INFORMATION:

APPLICANT: ST. GEORGE-HYSLOP, PETER H

APPLICANT: ROMMENS, JOANNA M

APPLICANT: FRASER, PAUL E

TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED

NUMBER OF SEQUENCES: 183

CORRESPONDENCE ADDRESS:

ADDRESSEE: TESTA, HURWITZ & THIBEAULT

STREET: High Street Tower - 125 High Street

CITY: Boston

STATE: Massachusetts

COUNTRY: U.S.A.

ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: FLOPPY disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

CURRENT APPLICATION DATA:

SOFTWARE: Patent Release #1.0, Version #1.30

APPLICATION NUMBER: US/08/967,101

FILING DATE: 10-NOV-1997

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/592,541

FILING DATE: 08-05-1995

ATTORNEY/AGENT INFORMATION:

NAME: Pitcher, Edmund R.

TELECOMMUNICATION INFORMATION:

TELEPHONE: (617) 248-7000

TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 137:

SEQUENCE CHARACTERISTICS:

LENGTH: 448 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: protein

US-08-967-101-137

Query Match 100.0%; DB 2: Length 448;

Best Local Similarity 100.0%; Pred. No. 1-231;

Matches 448; Conservative 0; Mismatches 0; Indels 0;

Gaps 0;

Query 1 MLTFMASDSEEEVCDERTSIMSAEPTPSQEGRPEDGENTAQWRSQENEEDGE

DP 60

Db 1 MLTFMASDSEEEVCDERTSIMSAEPTPSQEGRPEDGENTAQWRSQENEEDGE

DP 60

Query 61 DRYCSGVGRPPGLEEEELTKYAKHIVMLFVPTLCMVVVATKSYRFYTERNGQLI 120

Db 1 MLTFMASDSEEEVCDERTSIMSAEPTPSQEGRPEDGENTAQWRSQENEEDGE

DP 60

Query 61 DRYCSGVGRPPGLEEEELTKYAKHIVMLFVPTLCMVVVATKSYRFYTERNGQLI 120

Db 1 MLTFMASDSEEEVCDERTSIMSAEPTPSQEGRPEDGENTAQWRSQENEEDGE

DP 60

SUMMARIES

Result No.	Score	Query	Match	Length	DB	ID	Description
1	2336	100.0	448	2	US-08-967-101-137	Sequence 137, App	Sequence 137, App
2	2336	100.0	448	2	US-08-967-101-137	Sequence 137, App	Sequence 137, App
3	2320	99.3	447	2	US-08-875-972-29	Sequence 29, App	Sequence 29, App
4	1923	82.3	372	2	US-08-667-101-138	Sequence 138, App	Sequence 138, App
5	1923	82.3	372	2	US-08-592-541-138	Sequence 2, App	Sequence 2, App
6	1623	69.5	376	2	US-08-875-972-2	Sequence 13, App	Sequence 13, App
7	1467	62.8	467	2	US-08-967-101-134	Sequence 13, App	Sequence 13, App
8	1467	62.8	467	2	US-08-592-541-134	Sequence 2, App	Sequence 2, App
9	1467	62.8	467	3	US-08-670-964-2	Sequence 2, App	Sequence 2, App
10	1461	62.5	467	2	US-08-967-101-2	Sequence 2, App	Sequence 2, App
11	1461	62.5	467	2	US-08-592-541-2	Sequence 4, App	Sequence 4, App
12	1454	62.2	463	3	US-08-670-964-4	Sequence 4, App	Sequence 4, App
13	1438	61.6	407	2	US-08-875-972-4	Sequence 4, App	Sequence 4, App
14	1437	61.5	463	2	US-08-670-479-18	Sequence 18, App	Sequence 4, App
15	1418	60.7	467	2	US-08-967-101-4	Sequence 4, App	Sequence 4, App
16	1418	60.7	467	2	US-08-592-541-4	Sequence 166, App	Sequence 166, App
17	1150	49.2	541	2	US-08-967-101-166	Sequence 3, App	Sequence 3, App
18	1150	49.2	541	2	US-08-592-541-166	Sequence 3, App	Sequence 3, App
19	110	44.7	1294	2	US-08-819-288-3	Sequence 8, App	Sequence 8, App
20	108	4.6	1321	1	US-08-261-822A-3	Sequence 4, App	Sequence 4, App
21	108	4.6	1321	4	PCT-US95-07744A-3	Sequence 2, App	Sequence 2, App
22	101	4.3	1334	2	US-08-996-545-2	Sequence 8, App	Sequence 8, App
23	100	4.3	400	1	US-08-602-010A-8	Sequence 4, App	Sequence 4, App
24	95	4.1	391	1	US-07-816-283-4	Sequence 2, App	Sequence 2, App
25	95	4.1	391	1	US-08-417-103-4	Sequence 2, App	Sequence 2, App
26	93	4.0	391	1	US-07-815-283-2	Sequence 2, App	Sequence 2, App
27	93	4.0	391	1	US-08-417-103-2	Sequence 14, App	Sequence 6, App
28	93	4.0	391	1	US-08-417-103-14	Sequence 6, App	Sequence 1, App
29	93	4.0	509	2	US-08-031-392-6	Sequence 1, App	Sequence 1, App
30	91	3.9	400	1	US-08-602-010A-8	Sequence 8, App	Sequence 8, App
31	88.5	3.8	452	1	US-08-117-361C-2	Sequence 6, App	Sequence 6, App
32	88.5	3.8	3169	2	US-08-477-451-6	Sequence 3, App	Sequence 3, App
33	87	3.7	492	2	US-08-355-844-3	Sequence 3, App	Sequence 3, App
34	87	3.7	492	4	PCT-US95-16126-3	Sequence 3, App	Sequence 3, App

DESCRIPTION

US 08-875-972-29

Query Match Score 2310.5; DB 2; Length 447;
Best Local Similarity 99.3%; Pred. No. 4e-230; Matches 447; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 MLLFMASDSEEEVCDERTSLMSAESPRSCQGPGEDGENTAQWSQENEDGEEDP 60
Db 1 MLLFMASDSEEEVCDERTSLMSAESPRSCQGPGEDGENTAQWSQENEDGEEDP 60

QY 61 DRVCGSGVPGPGRGGLEEFLTKYGAHVIMLFYPTLGIVVATIKSVRFTEKNGOLI 120
Db 61 DRVCGSGVPGPGRGGLEEFLTKYGAHVIMLFYPTLGIVVATIKSVRFTEKNGOLI 120

QY 121 YTFETTEDPSYCGRLINSVNTLIMISVIVMFLVLYKCYCYKFTHGWLIMSSMLI 180
Db 121 YTFETTEDPSYCGRLINSVNTLIMISVIVMFLVLYKCYCYKFTHGWLIMSSMLI 180

QY 181 FIFTY TYGEVKLTYNVAMDYPFLTYWNFGAVGMCLHWKGPLVQAYLIMISALMA 240
Db 181 FIFTY TYGEVKLTYNVAMDYPFLTYWNFGAVGMCLHWKGPLVQAYLIMISALMA 240

QY 241 LVIKIYLPPEWSAWVILGAISVYDVLAVLCPKGPLRMLVETAQRNEPIFPALIYSSAMWV 300
Db 241 LVIKIYLPPEWSAWVILGAISVYDVLAVLCPKGPLRMLVETAQRNEPIFPALIYSSAMWV 300

QY 301 TYGMAKUDPSSQGALQPDPEMEEDSYDSFGPSYPVEFPEPLTGPGEELEEEERGV 360
Db 301 TYGMAKUDPSSQGALQPDPEMEEDSYDSFGPSYPVEFPEPLTGPGEELEEEERGV 359

QY 361 KLGIGDFIFYSVLVGKAATGSDWNTLACFTAILGLCILTLILAVEFKALPALPSI 420
Db 360 KLGIGDFIFYSVLVGKAATGSDWNTLACFTAILGLCILTLILAVEFKALPALPSI 419

QY 421 TFGLIFYFSTDNVRPFMDTIALSHOLYI 448
Db 420 TFGLIFYFSTDNVRPFMDTIALSHOLYI 447

RESULT 4
US 08-967-101-138
Sequence 138 Application US/08967101
Patent No. 584040

GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183
CORRESPONDENCE ADDRESS:
ADDRESS: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110

CURRENT APPLICATION DATA:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELECOMMUNICATION INFORMATION:

RESULT 5
US-08-592-541-138
Sequence 138 Application US/08592541
Patent No. 5986054

GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183
CORRESPONDENCE ADDRESS:
ADDRESS: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/967,101
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELECOMMUNICATION INFORMATION:

RESULT 138
US-08-592-541
Sequence 138 Application US/08592541
Patent No. 5986054

GENERAL INFORMATION:
APPLICANT: ST. GEORGE-HYSLOP, PETER H
APPLICANT: ROMMENS, JOHANNA M
APPLICANT: FRASER, PAUL E
TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED TO ALZHEIMER'S DISEASE
NUMBER OF SEQUENCES: 183
CORRESPONDENCE ADDRESS:
ADDRESS: TESTA, HURWITZ & THIBEAULT
STREET: High Street Tower - 125 High Street
CITY: Boston
STATE: Massachusetts
COUNTRY: U.S.A.
ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/592,541
FILING DATE: 10-NOV-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/592,541
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Pitcher, Edmund R.
TELECOMMUNICATION INFORMATION:

SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/967,101
 FILING DATE: 10-NOV-1997
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/592,541
 ATTORNEY/AGENT INFORMATION:
 NAME: Pitcher, Edmund R.
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (617) 248-7000
 TELEX: (617) 248-7100
 INFORMATION FOR SEQ ID NO: 134:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 467 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 US-08-967-101-134

Query Match 62.8%; Score 1467; DB 2; Length 467;
 Best Local Similarity 65.7%; Pred. No. 2e-142; Mismatches 39; Indels 42; Gaps 8;
 Matches 306; Conservative 39; US-08-967-101-134

Qy 24 ESPTPRSQCERGQPGDENTAAQRSQNEEDGEEDPDRYCSGVF----GRPPG---- 75
 Db 3 ELPAPLSYFQNAOMSNDHLSNTVSQNDNRERQEHINDR-SLGHEPEPLNSRGQNSRQ 61
 Qy 75 -----LEEPLTQXGAKHIVLMLFVPTLQCMIVVATISVRSRVEKQHQLIYTPTED 127
 Db 62 VEQDEEDEELTLYGAKHIVMLFPVILQMVVVATIKSVSFTRKGGLIYTPTED 121
 Qy 128 TPSVGORLINSVYNTLIMISVYNTMFLVYKRYCCKFTGHWLIMSSLMLLFETYIX 187
 Db 122 TETVGORALHSILNAIMISVYNTMFLVYKRYCCKFTGHWLISLSSLFFSFIX 181
 Qy 188 LGEVLKTYNAMDYPILLTWNFGAVGMYCIIHWKGPVLQAYLIMISALMALFIYL 247
 Db 182 LGEVFKTYNADVDTIVLWVQGMISIIHWKGPRLQQAYLIMISALMALFIYL 241
 Qy 248 PEWSAVVILGAISYDVAVLCPKGPLRMVYTAERNEPIFALIYSSAMWVYGMALK 307
 Db 242 PWTAMILAVIVTDLAVLCPKGPLRMVYTAQRNETIFPALIYSSTMWVLYMAEG 301
 Qy 308 DPSSQGAL-QLPYDPE-MEEDSYDSFGE---PSYPEVFEPPLTGPG----- 350
 Db 302 DPEAQRRVSKNSKNAESTERESQDVAENDDGGSEEEWAQRDSLGHPRSTPESRAAV 361
 Qy 350 EEL-----BEEEEGKVKGDFIFYSVLGVKAATGSDWNTLACFVAILGLCLT 402
 Db 362 QELSSSIAGEDPBERGVKLGLDEFIFYSVLGVKASATASDWNTIACFVAILGLCLT 421
 Qy 403 LLLAVYFKKALPALPISITFLPMDTFLASHQLYI 448
 Db 422 LLLAVYFKKALPALPISITFLPMDTFLASHQLYI 467

RESULT 8
 US-08-592-541-134
 Sequence 134, Application US/08592541
 Patent No. 5986054
 GENERAL INFORMATION:
 APPLICANT: ST. GEORGE-HYSLOP, PETER H
 APPLICANT: ROMMENS, JOHANNA M
 APPLICANT: FRASER, PAUL E
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 TITLE OF INVENTION: TO ALZHEIMER'S DISEASE
 NUMBER OF SEQUENCES: 183
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: TESTA, HURNITZ & THIBEAULT

RESULT 9
 US-08-570-964-2
 Sequence 2, Application US/08670964
 Patent No. 6010874
 GENERAL INFORMATION:
 APPLICANT: Hardy, John A.
 TITLE OF INVENTION: EARLY ONSET ALZHEIMER'S DISEASE

QY 308 DPSSQGAL-QLPYDPE-MEEDSYDSFGP---PSYPEVFEPPLTGYPG---- 350
 Db 302 DPAQRRTYSKNSYNAESTEREESODTVAENDGGFSEEMEAQDSDHGLPHRSPESSRAV 361
 QY 350 EEL-----EEPEERGTYKLGLGDFIFTYSVLYGKAATGSDPNTTLACFVALIGLCLT 402
 Db 362 QELSSSILAGEDDEERGTYKLGLGDFIFTYSVLYGAKASATASGDNTTIACFVALIGLCLT 421
 QY 403 LLLAVERFKALPAPLISITFGLLFYFSIDNLYRPFMDTFLASHQYI 448
 Db 422 LLLAIFKALPAPLISITFGLLFYFATDYLVCQFMDQLAFHQFYI 467

RESULT 11
 US-08-592-541-2
 Sequence 2, Application US/08592541
 Patent No. 5986054
 GENERAL INFORMATION:
 APPLICANT: ST. GEORGE-HYSLOP, PETER H
 APPLICANT: ROMMENS, JOHANNA M
 APPLICANT: FRASER, PAUL E
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 TO ALZHEIMER'S DISEASE
 NUMBER OF SEQUENCES: 183
 ADDRESSEE: TESPA, HURWITZ & THIBEAULT
 STREET: High Street Tower - 125 High Street
 CITY: Boston
 STATE: Massachusetts
 COUNTRY: U.S.A.
 ZIP: 02110
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patientin Release #1.0, version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/592,541
 FILING DATE:
 ATTORNEY/AGENT INFORMATION:
 NAME: Pitcher, Edmund R
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (617) 248-7000
 TELEFAX: (617) 248-7100
 INFORMATION FOR SEQ ID NO: 2:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 467 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 US-08-592-541-2

Db 182 LGEVRLTKYNAAMDPTLTLTVNNFGAYGRVCIHWKGPLVQQAYLIMISALMALVFIKYL 241
 QY 248 PEASAWVIGAISTYDLVAVLCPKSPRLMLVETAQERNEPFPLALYSAMWVTCGMALK 307
 Db 242 PEWTAWLAVISTYDLVAVLCPKSPRLMLVETAQERNEPFPLALYSIMWLYNMAEG 301
 Qy 308 DPSSQGAL-QLPYDPE-MEEDSYDSFGP---PSYPEVFEPPLTGYPG---- 350
 Db 302 DPEARRVSNSKNAESTERESQDQVAAANDGGSEEEWQDSDHLPHRSTPEERA V 361
 Qy 350 EEL-----EEPEERGTYKLGLGDFIFTYSVLYGKAATGSDPNTTLACFVALIGLCLT 402
 Db 362 QELSSSILAGEDDEERGTYKLGLGDFIFTYSVLYGAKASATASGDNTTIACFVALIGLCLT 421
 Qy 403 LLLAVERFKALPAPLISITFGLLFYFSIDNLYRPFMDTFLASHQYI 448
 Db 422 LLLAIFKALPAPLISITFGLLFYFATDYLVCQFMDQLAFHQFYI 467

RESULT 12
 US-08-670-964-4
 Sequence 4, Application US/08670964
 Patent No. 6010874
 GENERAL INFORMATION:
 APPLICANT: HARDY, JOHN A.
 TITLE OF INVENTION: EARLY ONSET ALZHEIMER'S DISEASE
 TITLE OF INVENTION: GENE AND GENE PRODUCTS
 NUMBER OF SEQUENCES: 4
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Smithline Beecham Corporation
 STREET: 709 Swedeland Road - UN2220; P.O. Box 15
 CITY: Philadelphia
 STATE: PA
 COUNTRY: USA
 ZIP: 19406
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: DOS
 SOFTWARE: FASTSEQ for Windows Version 2.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/670,964
 FILING DATE: 26-JUN-1996
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 60/001,142
 FILING DATE: 13-JUL-1995
 APPLICATION NUMBER: 60/001,501
 FILING DATE: 18-JUL-1995
 ATTORNEY/AGENT INFORMATION:
 NAME: Han, William T
 REGISTRATION NUMBER: 34,344
 REFERENCE/DOCKET NUMBER: P-0358
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 610-270-5219
 TELEFAX: 610-270-5090
 TELEX:
 INFORMATION FOR SEQ ID NO: 4:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 463 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 US-08-670-964-4

Query Match 62.5%; Score 1451; DB 2; Length 467;
 Best Local Similarity 65.5%; Pred. No. 8.1e-142; Indels 42; Gaps 8;
 Matches 305; Conservative 39; Mismatches 80; Indels 42; Gaps 8;

Qy 24 ESSTPRSQEGRGPGEDBENTAWRSQNEEDBEDPDRYVCSCVP----GRPPG---- 75
 Db 3 ELAPLSYFQNACMDSDHLSNTVRSDQNDNR-SGHPPELSNGPQGNRQ 61
 Qy 75 -----LEELLTLYKYGAKHVINLFPVYPTLCLMIVVATIKSVRFYETEKNGOLYTPTFED 127
 Db 62 VVQDDEEDEELLTLYKYGAKHVINLFPVYPTLCLMIVVATIKSVSFYTRKGOLYTPTFED 121
 Qy 128 TPSVQORLNSVNTLIMISVTPATIPLWLVLYKRCXKFIRGWLIMSLMLFLFTY 187
 Db 122 TEVGVORALHSILNAAIMTSVIVMTILVVLVLYKRCVKVIAWLISSLJLFFSFY 181
 Qy 188 LGEVRLTKYNAAMDPTLTLTVNNFGAYGRVCIHWKGPLVQQAYLIMISALMALVFIKYL 247

Query Match 62.3%; Score 1454; DB 3; Length 463;
 Best Local Similarity 65.3%; Pred. No. 4.2e-141;
 Matches 303; Conservative 40; Mismatches 79; Indels 42; Gaps 8;

Qy 24 ESPTPRSCQEGRGPGEDGENTAQWSQEENEDGEEDPDRYVCSCVP--GRPPG---- 75

REFERENCE/DOCKET NUMBER: P50361
 TELEPHONE: 610-270-5219
 TELEFAX: 610-270-5090
 TELEX:
 INFORMATION FOR SEQ ID NO: 18:
 SEQUENCE CHARACTERISTICS:
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: peptide
 HYPOTHETICAL: NO
 FRAGMENT TYPE: N-terminal
 ORIGINAL SOURCE:
 US -08-670-479-18

Query Match Score 1437; DB 2; Length 463;
 Best Local Similarity 64.9%; Pred No 2, 3e-139; Gaps 8;
 Matches 301; Conservative 41; Mismatches 80; Indels 42; Gaps 8;

QY 24 ESPPRSRSCOEGRQGPEDGENTAQWRSQEEDGEEDPDRYVCSCGV--GRPPG-----
 Db 3 ELAPLSTYQNAQMSNDHLS--NTNDNRERQHNDRSLRGHEPLNSGRQNSRQVV 59

QY 75 ---LEEEELTKYGAHKVIMFLPVPTLQMVVATIKSVRFTEKGQLIYTPPTEDP 129
 Db 60 EQDDEEDEEELTKYGAHKVIMFLPVPTLCMVVYVATIKSVRFTEKRDGLIYTPPTEDT 119

QY 130 SVOQRNLNSVLTNLMISVTVMMTFLVLYKTRCYKFTHGWLMSSMLFLFTYIYLG 189
 Db 120 TVGORALHSILNAIMISVTVMMTFLVLYKTRCYKVTHAWLIISSLLFFESFYLG 179

QY 190 EVLKTYNVAMDYPTLTLTYYNFGAAGMYCITHWKGPVLQQAYLIMISALMAYFIKYLPE 249
 Db 180 EVLKTYNVADYTTVALIWNFGTVGMISIHWKGPRLQQAYLIMISALMAYFIKYLPE 239

QY 250 WSAVWILGAISVTDLVAVLCPKGPLRMLVETAQRNEPFPALIYSSAMWTVGMAKLD 309
 Db 240 WTAWILAVISVQDVLVAVLCPKGPLRMLVETAQRDETLPALIYSSTNWLVNMAEGP 299

Qy 310 SSQGAL---OLPYDPE-MEEDSYDFGE --PSYBEVFPEPLTGPGP-----EE 351
 Db 300 EAQRVSKSKYNAESTERESQDVAENDGGFSEWEAQDSILGPHTSPPERAYOE 359

QY 352 L-----EEEEERGVKLGLGDFIFSYLVGKAATGSDWNTLACFVAILIGCLTL 404
 Db 360 LSSTSLAGEDPERGVKLGLGDFIFSYLVGKAATASGSDWNTLACFVAILIGCLTL 419

QY 405 LLAVFKKALPALPISITGLIFPFSTDNLYRPMDTЛАSHOLYI 448
 Db 420 LLAVFKKALPALPISITGLIFPFSTDNLYRPMDTЛАSHOLYI 463

RESULT 15
 US-08-967-101-4
 Patent No. 5840510
 GENERAL INFORMATION:
 APPLICANT: ST. GEORGE-HYSLOP, PETER H
 APPLICANT: ROMMENS, JOHANNA M
 TITLE OF INVENTION: GENETIC SEQUENCES AND PROTEINS RELATED
 NUMBER OF SEQUENCES: 183
 CORRESPONDENCE ADDRESS:
 ADDRESS: TESTA, HIRWITZ & THIBEAUT
 STREET: High Street Tower - 125 High Street
 CITY: Boston
 STATE: Massachusetts
 COUNTRY: U.S.A.

; ZIP: 02110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/967,101
; FILING DATE: 10-NOV-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/592,541
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Pitcher, Edmund R.
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 248-7000
; TELEFAX: (617) 248-7100
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 467 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-967-101-4

Query Match Score 1418; DB 2; Length 467;
 Best Local Similarity 61.9%; Pred No. 2.1e-137;
 Matches 289; Conservative 48; Mismatches 86; Indels 44; Gaps 6;

QY 24 ESPPRSRSCOEGRQGPEDGENTAQWRSQEEDGEEDPDRYVCSCGVP---GRPPG---- 75
 Db 3 EIPAPLSYQNAQMSNDHLS--NTNDNRERQHNDRSLRGHEPLNSGRQNSRQVV 62

Qy 75 -----LEEEELTKYGAHKVIMFLPVPTLQMVVATIKSVRFTEKGQLIYTPPTEDT 128
 Db 63 VEQDEEEDEEELTKYGAHKVIMFLPVTLQMVVATIKSVRFTEKRDGLIYTPPTEDT 122

QY 129 PSVGORLNSVLTNLMISVTVMMTFLVLYKTRCYKFTHGWLMSSMLFLFTYIYL 188
 Db 123 ETVGORALHSILNAIMISVTVMMTFLVLYKTRCYKVTHAWLIISSLLFFESFYL 182

QY 189 GEVLTKYNVAMDYPTLTLTYYNFGAAGMYCITHWKGPVLQQAYLIMISALMAYFIKYLPE 248
 Db 183 GEVFKTYNVADYTTVALIWNFGTVGMISIHWKGPRLQQAYLIMISALMAYFIKYLPE 242

QY 249 EWSANVILGAIISVTDLVAVLCPKGPLRMLVETAQRNEPFPALIYSSAMWTVGMAKLD 308
 Db 243 EWTANLILATISVQDVLVAVLCPKGPLRMLVETAQRNETLPALIYSSTNWLVNNAEGD 302

QY 309 PSSQGALQLPYDPME-----EDSYDSFGEPSYPEVFPPLTGPGP----- 350
 Db 303 PEAQ --RRVPKNPKNTQRERETDGSGNDDGGFSEWAQRDSSHGPFRSTPESRAA 360

QY 350 -EEL-----EEEEERGVKLGLGDFIFSYLVGKAATGSDWNTLACFVAILIGCL 401
 Db 361 VQELSGSILTSEDPBPGRVNLGLGDFIFSYLVGRASATASGDWNTLACFVAILIGCL 420

QY 402 TLILLIAVEKALPALPISITGLIFPFSTDNLYRPMDTЛАSHOLYI 448
 Db 421 XLLILIAVKKXPAPISITGLIFPFSTDNLYRPMDTЛАSHOLYI 467

Search completed: March 18, 2000, 19:55:31
 Job time: 3229 sec

Copyright (c) 1993 - 1998 Compugen Ltd.	Gencore version 4.5	36	98	4.2	461	2	T11829
OM protein - protein search, using sw model.		37	98	4.2	1836	2	JS0648
Run on:	March 18, 2000, 14:03:43 ; Search time 41.25 Seconds 512.287 Million cell updates/sec	38	98	4.2	1836	2	I51964
Title:	US-08-509-359B-137	39	98	4.2	1836	2	I51964
Perfect score:	2336	40	98	4.2	1836	2	I64893
Sequence:	1 MLTFMASDSPEEVCDERTSL.....STDNLVRPFMDTLASHOLYI 448	41	98	4.2	1835	2	I54323
Scoring table:	BLOSUM62	42	98	4.2	217	2	S01095
Searched:	142080 seqs, 47169319 residues	43	97.5	4.2	1695	2	JE0084
Database :	PIR_62; *	44	97	4.2	441	2	S13425
Word size :	0	45	97	4.2	381	2	T11440
Number of hits that pass the threshold :	1420808				447	2	S52968
1: pirl;*							
2: pir2;*							
3: pir3;*							
4: pir4;*							
Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.							
SUMMARIES							
Result No.	Score	Query Match	Length	DB ID	Description		
- - -	- - -	- - -	- - -	- - -	- - -	- - -	- - -
1	2316	100.0	448	2	I58098	E5.1 Protein - human	RESULT 1
2	2338	99.7	448	2	A56993	C;Species: Homo sapiens (man)	
3	2215	94.8	442	2	J39174	C;Accession: I58098	
4	1624	69.5	449	2	JC5391	R; Rogaev, E.I.; Sherrington, R.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Liang, Y.; Cohen, D.; Lannfelt, L.; Fraser, P.E.; Rommens, J.M.; St George-Hyslop, P.H.	
5	1468	62.8	467	2	I78388	Nature 376, 775-778, 1995	
6	1457	62.8	467	2	S58336	A;Title: Familial Alzheimer's disease in kindreds with missense mutations in a gene on chromosome 14.	
7	1454	62.2	463	2	S63683	A;Reference number: I58098;	
8	1449	62.1	433	2	JC5390	A;Accession: I58098	
9	1447	61.9	467	2	JC5050	A;Molecule type: mRNA	
10	1440	61.6	463	2	JC5081	A;Residues: 1-448 <RES>	
11	1035	44.3	374	2	S63684	A;Cross-references: GB:L44577; PID:9950347; NID:9950348	
12	967	41.4	461	2	S60253	C;Genetics:	
13	524.5	22.5	358	2	T15184	A;Gene: 95-1	
14	513	22.0	453	2	T00724	C;Superfamily: Presenilin	
15	274	11.7	465	2	A43459		
16	119	5.1	531	2	T11596		
17	110	4.7	2016	2	A38195		
18	109	4.7	1840	1	CHRT01		
19	106	4.5	398	2	H75043		
20	104	4.5	1476	1	A39901		
21	104	4.5	1476	1	A40303		
22	104	4.5	826	2	T02268		
23	103.5	4.4	1450	2	JC6139		
24	103	4.4	382	2	S47882		
25	102.5	4.4	893	2	A47550		
26	102	4.4	379	2	I48135		
27	101.5	4.3	1681	2	S00320		
28	100.5	4.3	1951	2	A60054		
29	100.5	4.3	1983	2	T01900		
30	100.5	4.3	766	2	S36646		
31	100.5	4.3	324	2	E70488		
32	100	4.3	592	2	JC6178		
33	99.5	4.3	379	2	A33995		
34	99	4.2	2019	2	521		
35	99	4.2	521	2	T11710		

ALIGNMENTS

RESULT 1						
Query	MLTFMASDSPEEVCDERTSL.....STDNLVRPFMDTLASHOLYI 448	Match	100.0%	Score	2336;	DB 2;
Best		Local	100.0%	Pred.	No. 8.1e-167;	Length 448;
Matches		Conservative	0;	Mismatches	0;	Gaps 0;
QY	MLTFMASDSEEVCDERTSLMSAESPPTPSQCOEGRQPEDGEDEDP 60	Db	1	MLTFMASDSEEVCDERTSLMSAESPPTPSQCOEGRQPEDGEDEDP 60	1	MLTFMASDSEEVCDERTSLMSAESPPTPSQCOEGRQPEDGEDEDP 60
YQ	MLTFMASDSEEVCDERTSLMSAESPPTPSQCOEGRQPEDGEDEDP 60	Qy	61	DRYVCSGVGRPQEELITLKYGAKHVNMLFVVTLCMVIVVATIKSRYEFTENGQLI 120	61	DRYVCSGVGRPQEELITLKYGAKHVNMLFVVTLCMVIVVATIKSRYEFTENGQLI 120
Db	MLTFMASDSEEVCDERTSLMSAESPPTPSQCOEGRQPEDGEDEDP 60	Db	61	DRYVCSGVGRPQEELITLKYGAKHVNMLFVVTLCMVIVVATIKSRYEFTENGQLI 120	61	DRYVCSGVGRPQEELITLKYGAKHVNMLFVVTLCMVIVVATIKSRYEFTENGQLI 120
Qy	YTPTEDTPSVGQRQLNSYNTLIMISVYVMTIFLVVLYKRYCKFIFIQWLIMSSLMLI 180	Qy	121	YTPTEDTPSVGQRQLNSYNTLIMISVYVMTIFLVVLYKRYCKFIFIQWLIMSSLMLI 180	121	YTPTEDTPSVGQRQLNSYNTLIMISVYVMTIFLVVLYKRYCKFIFIQWLIMSSLMLI 180
Db	YTPTEDTPSVGQRQLNSYNTLIMISVYVMTIFLVVLYKRYCKFIFIQWLIMSSLMLI 180	Db	121	YTPTEDTPSVGQRQLNSYNTLIMISVYVMTIFLVVLYKRYCKFIFIQWLIMSSLMLI 180	121	YTPTEDTPSVGQRQLNSYNTLIMISVYVMTIFLVVLYKRYCKFIFIQWLIMSSLMLI 180
Qy	FIFTTYIYLGEVLKTYNVANDYPTLTVYNEGAVGMVCIHWKGPLVLYQDAYLIMISALMA 240	Qy	181	FIFTTYIYLGEVLKTYNVANDYPTLTVYNEGAVGMVCIHWKGPLVLYQDAYLIMISALMA 240	181	FIFTTYIYLGEVLKTYNVANDYPTLTVYNEGAVGMVCIHWKGPLVLYQDAYLIMISALMA 240
Db	FIFTTYIYLGEVLKTYNVANDYPTLTVYNEGAVGMVCIHWKGPLVLYQDAYLIMISALMA 240	Db	181	FIFTTYIYLGEVLKTYNVANDYPTLTVYNEGAVGMVCIHWKGPLVLYQDAYLIMISALMA 240	181	FIFTTYIYLGEVLKTYNVANDYPTLTVYNEGAVGMVCIHWKGPLVLYQDAYLIMISALMA 240
Qy	LVFKYLPLPSAWTILGAISVQDNLVAVLQPKGLPMLVTAQERNEPFLVYSSAMW 300	Qy	241	LVFKYLPLPSAWTILGAISVQDNLVAVLQPKGLPMLVTAQERNEPFLVYSSAMW 300	241	LVFKYLPLPSAWTILGAISVQDNLVAVLQPKGLPMLVTAQERNEPFLVYSSAMW 300
Db	LVFKYLPLPSAWTILGAISVQDNLVAVLQPKGLPMLVTAQERNEPFLVYSSAMW 300	Db	241	LVFKYLPLPSAWTILGAISVQDNLVAVLQPKGLPMLVTAQERNEPFLVYSSAMW 300	241	LVFKYLPLPSAWTILGAISVQDNLVAVLQPKGLPMLVTAQERNEPFLVYSSAMW 300
Qy	TVMGAKLDPSSQGQLPQDPMEMEDSYSDFGEPSEPSYEFPPPLTGYPGFEELEEEERYV 360	Qy	301	TVMGAKLDPSSQGQLPQDPMEMEDSYSDFGEPSEPSYEFPPPLTGYPGFEELEEEERYV 360	301	TVMGAKLDPSSQGQLPQDPMEMEDSYSDFGEPSEPSYEFPPPLTGYPGFEELEEEERYV 360
Db	TVMGAKLDPSSQGQLPQDPMEMEDSYSDFGEPSEPSYEFPPPLTGYPGFEELEEEERYV 360	Db	301	TVMGAKLDPSSQGQLPQDPMEMEDSYSDFGEPSEPSYEFPPPLTGYPGFEELEEEERYV 360	301	TVMGAKLDPSSQGQLPQDPMEMEDSYSDFGEPSEPSYEFPPPLTGYPGFEELEEEERYV 360
Qy	KLGIGDFITYFSVLYKGKAATGSGDWNTTLACFVAILGICLTLIAYFKPALPISI 420	Qy	361	KLGIGDFITYFSVLYKGKAATGSGDWNTTLACFVAILGICLTLIAYFKPALPISI 420	361	KLGIGDFITYFSVLYKGKAATGSGDWNTTLACFVAILGICLTLIAYFKPALPISI 420
Db	KLGIGDFITYFSVLYKGKAATGSGDWNTTLACFVAILGICLTLIAYFKPALPISI 420	Db	361	KLGIGDFITYFSVLYKGKAATGSGDWNTTLACFVAILGICLTLIAYFKPALPISI 420	361	KLGIGDFITYFSVLYKGKAATGSGDWNTTLACFVAILGICLTLIAYFKPALPISI 420
Qy	TFLGLIFYFSTDNLVRFMFLASHOLYI 448	Qy	421	TFLGLIFYFSTDNLVRFMFLASHOLYI 448	421	TFLGLIFYFSTDNLVRFMFLASHOLYI 448
Db	TFLGLIFYFSTDNLVRFMFLASHOLYI 448	Db	421	TFLGLIFYFSTDNLVRFMFLASHOLYI 448	421	TFLGLIFYFSTDNLVRFMFLASHOLYI 448

QY	7	SDSEDEVCDERTSLMSAESITPRSQEGRGPDGEDNTAQRSGNEDBBDPDRYVC S 66	QY	249	EWSAWVILGAISSVYDVLAVLCPKGPLRMLVYETAQERNEPITPPLIYSSAMWVTVGMAKLD 308
Db	5	SDSEDECNRSLTISSESPPLPSQDGYSYRERQPISTQNED----- 58	Db	243	EWTAWLALIVSITVLAVLCPKGPLRMLVYETAQERNETLPALIYSTMWLVNMAEGD 302
QY	67	GVP-GRPPGLI-----EPEELTKYGAHKYIMLFPTVLCMIVYYATIKSYRFYT 113	QY	309	PSSQGALQLPYDPEME-----EDSYDSFGEPSSPEVFVPEPLTGYP----- 350
Db	58	-VPNGRTISGDAYNSETTVNEEEETLKGARHYIMLFPTVLCMIVYYATIKSYFT 116	Db	303	PEAQ - .RVRPKNPKYNTORAERETDPSGSNDGGSEEEAQRDSHGPFRSTPESRAA 360
QY	114	EKGQLIYTFETDTPSGORLINSVNTLIMISVWMTFLFLVLYKRYCKFKIIGWL I 173	QY	350	-EEL-----EEEERGVKLJGLDEFIFYSVLGKAATGSGDWNNTLACFVAILIGLCL 401
Db	117	EKGQLIYTFETDTPSGERLINSVNTLIMISVWMTFLFLVLYKRYCKFKIIGWL I 176	Db	361	VOELSGSILTSEDPPERGVKLJGLDEFIFYSVLGKAATASGDDWNTIACFVAILIGLCL 420
QY	174	MSSMLLFLFTTYIYGEVLKTYNYAMDYPILLTYWNFGAVGMYCITHKGKPLVQOAYLI 233	QY	402	TLLILAYFEKKALPALPISITFGLLIFYFSTDNDLVRPPMDTLASHQLYI 448
Db	177	LSSMLLFLFTTYIYSEFRTYNIAMDYPLEMVWNFGAVGMICITHKGKPLVQOAYLI 236	Db	421	TLLILAYFEKKALPALPISITFGLLIFYFATDYLVQPPMDOLAFHQFYI 467
QY	234	MISALMALVFIKYLPDEWSAWVILGAISSVYDVLAVLCPKGPLRMLVYETAQERNEPITPPLI 293	RESULT 6		
Db	237	MISALMALVFIKYLPDEWSAWVILGAISSVYDVLAVLCPKGPLRMLVYETAQERNEPITPPLI 296	558396		
QY	294	YSSAMWVYGMALDKPSSQAL---OLP-----DPMEEDSYDSFGEPSPYEVFEPLTG Y 347			
Db	297	YSSAMWVYGMAD-SATADGMRMNOQVHIDRNTPAGANSVTEAAETRQ-----QS 348			
QY	348	PGEELEEEERGVKLJGLDEFIFYSVLGKAATGSGDWNNTLACFVAILIGLCLTLIA 407			
Db	349	NLSSSDPDERGVKLJGLDEFIFYSVLGKAATASGDDWNNTLACFVAILIGLCLTLIA 408			
QY	408	VFKKALPALPISITFGLLIFYFSTDNDLVRPPMDTLASHQYI 448			
Db	409	VFKKALPALPISITFGLLIFYFSTDNDLVRPPMDTLASHQYI 449			
QY	5	RESULTS 178388			
C	Species: Mus musculus (house mouse)				
C	Date: 27-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 29-Sep-1999				
C	Accession: I78388				
R	Sherrington, R.; Rogaeva, E.I.; Liang, Y.; Rogaeva, E.A.; Levesque, G.; Ikeda, M.; Chi, E.; Rommens, J.M.; St. George-Hyslop, P.H.				
D	Nature 375, 754-760, 1995				
A	Title: Cloning of a gene bearing missense mutations in early-onset familial Alzheimer				
A	Reference number: 158095; MUID:95319502				
A	Accession: S58396				
A	Molecule type: mRNA				
A	Residues: 1-467 <SHE>				
A	Cross-references: EMBL:142110; NID:9904118; PID:9904119				
R	Viadal, R.; Ghiso, J.; Wisniewski, T.; Frangione, B.				
F	FEBS Lett. 393, 19-23, 1996				
A	Title: Alzheimer's presenilin 1 gene expression in platelets and megakaryocytes.				
A	Reference number: S71401; MUID:96397521				
A	Accession: S71401				
A	Status: not compared with conceptual translation				
A	Molecule type: mRNA				
A	Residues: 24-32/254-256,290-292;316-317,376-379 <VIEW>				
A	Alternative source: Dani megakaryotic cell line (ATCC CRL-9792) and platelets				
C	Genetics:				
A	Gene: GDB:PSEN1; AD3; PAD; S182; PS1				
A	Cross-references: GDB:135682; OMIM:104311				
A	Map position: 14q24.3-14q24.3				
C	Superfamily: presenilin				
C	Keywords: alternative splicing; Alzheimer's disease; glycoprotein; transmembrane protein				
F	B2/100/Domain: transmembrane #status predicted <TM1>				
F	113/154/Domain: transmembrane #status predicted <TM2>				
F	16/185/Domain: transmembrane #status predicted <TM3>				
F	19/213/Domain: transmembrane #status predicted <TM4>				
F	221-238/Domain: transmembrane #status predicted <TM5>				
F	244-264/Domain: transmembrane #status predicted <TM6>				
F	281-301/Domain: transmembrane #status predicted <TM7>				
F	408-428/Domain: transmembrane #status predicted <TM8>				
F	433-453/Domain: transmembrane #status predicted <TM9>				
F	279-405/Binding site: carbohydrate (Asn) (covalent)				
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Best Local Similarity	64.0%; Pred. No. 3.9e-10;				
Matches	299; Conservative 46; Mismatches 78; Indels 44; Gaps 6;				
QY	24	ESPPIRSCQEGRQGPEDGNTAQWQSQEENEDBPPDRYVCSGYP----GRPPG---- 75			
Db	3	EIPAPLSYFQNAQMSEDSHSSAIRSQNDQSERQQHDRLDNPPISINGRPOSNSRQ 62			
QY	75	-----LEPEELTKYGAHKYIMLFPTVLCMIVYYATIKSYRFYETEKNGQJYTPPTEDT 128			
Db	63	VEQDSEEDEELTKYGAHKYIMLFPTVLCMIVYYATIKSYFPTRKDGQJYTPPTEDT 122			
QY	24	ESPTRSCQEGRQGPEDGNTAQWQSQEENEDBPPDRYVCSGYP----GRPPG---- 75			
Db	3	ELPAPLSYFQNAQMSEDDHNLNTVSQNDRERQERANDRR-SLGHPPELNSGRQ 61			
QY	75	-----LEPEELTKYGAHKYIMLFPTVLCMIVYYATIKSYRFYETEKNGQJYTPPTEDT 127			
Db	62	VVEQQDSEEDELTKYGAHKYIMLFPTVLCMIVYYATIKSYRFYETRDQGLYTPPTEDT 122			
QY	129	PSVGORLINSVNTLIMISVWMTFLFLVLYKRYCKFKIIGWLIMSLMFLFTYIY 188			
Db	123	ETVYGRALISLNAIMSVIVINTLFLVLYKRYCKKVHAWLISLLLFFSIYL 182			
QY	189	GEVLRKTYNAVMDYPILLTYWNFGAVGMVCIHWKGKPLVLOQAYLIMISALMVFYKL P 248			
Db	183	GEVFKTYNAVDYTVALLWNFGVGMIAIHKGKPLVLOQAYLIMISALMVFYKL P 242			

presenillin 1 protein isoform 467 - lesser mouse lemур						
C;Species: Microcebus murinus (lesser mouse lemур)	C;Accession: JC5080	C;Date: 31-Jan-1997 #sequence_revision 31-Jan-1997	C;Text_Change 29-Sep-1999			
R;Bioclim: Biophys. Res. Commun. N.; Czech, C.; Pradier, L.; Petter, A.; Bons, N.; Bellis, A.; Mestre-Frances, N.; Mestre-Frances, N.; Czech, C.; Pradier, L.; Petter, A.; Bons, N.; Bellis, A.; Title: Molecular cloning, sequencing, and brain expression of the presenillin 1 gene in						
A;Reference number: JC5080; MJD:97079199						
A;Accession: JC5080						
A;Status: nucleic acid sequence not shown						
A;Gene: ps1						
A;Map position: 14						
C;Superfamily: presenillin						
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F;133-154/Domain: transmembrane #status predicted <TM2>						
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F;211-236/Domain: transmembrane #status predicted <TM5>						
F;244-261/Domain: transmembrane #status predicted <TM6>						
F;248-268/Domain: transmembrane #status predicted <TM7>						
Query Match Similarity 61.9%; Score 1447; DB 2; Length 467;						
Best Local Similarity 64.4%; Pred. No. 1.4e-100;						
Matches 300; Conservative 41; Mismatches 83; Indels 42; Gaps 7;						
QY 24 ESSPPRSCQEGRQPEDGENTAQWRSQEENDGEEDPDRYTCGVP -----GRPPG---- 75						
Db 3 ELPAPLSPYQDQNAQMSDNDHNLVQDQNSDNTYQSDNDHNLVQDQNSDNTYQSDNDHNLVQDQNSGP 61						
QY 75 -----LEBELTLYKGAHKVIMLFVPTVLCMIVVATIKSVRETYKETKNGQOLIYTPTED 127						
Ddb 62 VVERDEEEDEBLLTLYKGAHKVIMLFVPTVLCMIVVATIKSVFSTRKDGQLIYTPTED 121						
QY 128 TPSVGORLJNSVLNLIMISVIVVVTCIILVLYKKYCYKEFHGWLMISLMULFLTYIX 187						
Ddb 122 TETVQRALHSJSLNAAIMSVIVVVTCIILVLYKKYCYKVIAHAWLISLLUFFFFSTY 181						
QY 188 LGEVLTKNYAMDPYLTLTWNFGAYGMCIHWKGPLVLQOAYLIMISALMAVLFKYL 247						
Ddb 182 LGEFVKTYNNYAVDITVLLWNFGVYGMISHWKWPLRQOAYLIMISALMAVLFKYL 241						
QY 248 PEWSAWVILGAISYDVLAVLCPKGPLRMVYTAERNEIPFPALIYSSAMWTVGMAKL 307						
Ddb 242 PEWTAWLILAVISYDVLAVLCPKGPLRMVYTAERNEIPFPALIYSSAMWTVGMAKL 301						
QY 308 DPSSQGAL--QLPD-----PEMEEDSYDSFGESPYEFP----PLTGYPG 349						
Ddb 302 DPEAQRRYSRNTKINAQGTREQAQSVPENDGGFSEWEAQDOSLGPHRSTSVAARV 361						
QY 350 EEL-----EEEEEERGVKGVLGDFIFTYFSLVKGAAATGSDWNITLACFVAILIGCLT 402						
Ddb 362 QEISSSPASDPEERGVKGVLGDFIFTYFSLVKGAAATGSDWNITLACFVAILIGCLT 421						
QY 403 LLILAVFKKALPALPISITGLIYESTDNLVRPFMDTLASHOLYI 448						
Ddb 422 LLILIAFKKALPALPISITGLVFTATDVLQPFMDLAFHOFYI 467						

Biochem. Biophys. Res. Commun. 228, 430-439, 1996
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A Reference number: JC5080. MUD:9709199

A;Content: brain
A;Accession: JC5081
A;Status: nucleic acid sequence not shown
A;Molecule type: mRNA
A;Residues: 1-465 <CAI>
A;Cross-references: EMBL:Z11333
C;Comment: This protein is an intermembrane protein with seven transmembrane domains.

C; Genetics;
A; Gene: ps1
A; Map position: 14
C; Superfamily: presenilin

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C: Keywords: transmembrane Protein
C: 78-96/Domain: transmembrane #status predicted <TM1>
F: 129-150/Domain: transmembrane #status predicted <TM2>
F: 160-181/Domain: transmembrane #status predicted <TM3>
F: 191-209/Domain: transmembrane #status predicted <TM4>
F: 217-234/Domain: transmembrane #status predicted <TM5>
F: 220-277/Domain: transmembrane #status predicted <TM6>
F: 404-424/Domain: transmembrane #status predicted <TM7>

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 Best Local Similarity 64.28%; Pred. No. 4.7e-100;
 Matches 300; Conservative 39; Mismatches 80; Indels 48; Gaps 8;

Qy	75	- - - - - L E E E E T L K Y G A K H Y I M L F V P V T L C M V V A T I K S V R F Y T E K N G O L I T Y P F T E
Db	57	P V V E R D E E D E E T L K Y G A K H Y I M L F V P V T L C M V V A T I K S V S P Y T R K D G O L I T Y P F T E
	116	

Qy	127	DTPSVGORLNSVNTLIMISIVMFTFLVWLYKRCYKFHGWLMSSMLLFIFTYI	186
	117	DTETVGRALHSVYNAIMISIVMFTFLVWLYKRCYKVIAHWLIISSLLFFFSFI	176
Db			

	Qy	24 / LPFWSWAVLILGASVIVDYLAVVLCPKGIRMLVTAQERNLFLPALIYSSTMMWLVNMAE 306
	Db	237 LPENTWAVLILAVSYDYLAVVLCPKGIRMLVTAQERNLFLPALIYSSTMMWLVNMAE 296
Ov	307 LDPPSSGAL-OPLYD-----PEMEEDSYDSGEPSYEPFEP----PLTGYP 348	

Qy	349	GEEL	-----	-REEEERGVKLJGDFIYFVNLGKAATGSDWNTLACFVAILGLCL	401
Db	297	GDPEAQRVSNTKNTNAOGTEREAQASYPENDDGFSSEMEAORDSQLPHRSTSVA	356		

Qy	402	TLLLIAVFKKALPALPISITGLIFYFSTDNLRYRPEMDTLASHQYI	448
Db	357	VQEISSIPASEDPEEGVKGJGDFVFSVLYGKASATASGDNTTICFVAILLGCL	416

RESULT 11
Db 417 TLLLAFFKALPALPISITFGLVYFYATDYLVQPFMDQLAFHOFYI 463

S63684 Presenilin 1, splice form 374 - human
N:Alternative name: Alzheimer's disease protein 3
C:Species: *Homo sapiens* (human)

C;Date:20-Jul-1996 #sequence_revision 13-Mar-1997 #text_change 29-Sep-1999
 C;Accession: S63884
 R;Sahara, N.;Yanagi, Y.;Takagi, H.;Kondo, T.;Okochi, M.;Usami, M.;Shirasawa, T.
 FEBS Lett. 381:7-11, 1996

A; Title: Identification and characterization of presenilin 1-46/, 1-463 and 1-34/.
A; Reference number: S63683; MUID:96193901
A; Accession: S63684

A;Title: Mutation of a putative sperm membrane protein in *Caenorhabditis elegans* prev
A;Reference number: A43459; MUID:2407340
A;Status: Preliminary; not compared with conceptual translation
A;Accession: A43459
A;Molecule type: DNA; mRNA
A;Residues: 1-465 <LHE>
A;Cross-references: EMBL:214067; NID:96868; PID:96869; EMBL:714066; NID:96870; PID:96
A;Experimental source: strain Bristol N2
A;Note: the nucleotide sequence was submitted to the EMBL Data Library, July 1992
C;Genetics:
A;Introns: 69/3; 154/3; 200/1; 224/3; 300/1; 386/1; 435/1
A;Description: Genomic sequence for *Arabidopsis thaliana* BAC F22O13.
A;Number: Z14200
A;Accession: T0074
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 1-453 <SH>>
A;Cross-references: EMBL:AC003981; NID:93063438; PID:93063457; GSPDB:GN0059; ATSP:F22O13
C;Genetics:
A;Gene: ATSP:F22O13.19
A;Position: 1
A;Introns: 108/1
C;Superfamily: presenilin

Query Match 11.7%; Score 274; DB 2; Length 465;
Best Local Similarity 21.5%; Pred. No. 3.4e+13; Gaps 14;
Matches 93; Conservative 80; Mismatches 127; Indels 132; Gaps 14;

QY 114 EKGOLIYPPFTEDT-PSVGQRLINNSVLN---TLIMISVIVMTIFLYKRYCYKF 167
Db 42 EVNSLSSKTYFLDPSFEEQTGNLILDFENGVTIVLGCVSFIMAF--VLEDFER-RI 97

QY 168 IRGWLIIMSIIMLLF-----LEFTYTYGEVLTYKTNVAMDYPTILL----TWNFGA 213
Db 98 VKAWLITLSCCLLIFGVSAQTLDHDMFSQVTDODNNQY---YMTIVLIVVPTVYGF- 152

QY 214 VGMTCIHWKGPLVLOQAYIJMISALMAYEIKLPESWAIVLGAISVYDLIVLCPKG 273
Db 152 -GIAFFSSNLSLHHQIFVVTNCISLISVYLRVFSKTWTFLWFLDFAVLAPMGP 210

Query Match 22.0%; Score 511; DB 2; Length 453;
Best Local Similarity 31.1%; Pred. No. 5.2e+31; Gaps 12;
Matches 136; Conservative 78; Mismatches 130; Indels 94; Gaps 12;

QY 88 VIMLFVPTVLCKMIVVVVATIKSYRFETYTKN---GQLITYPTFTDDTPSYVGQRLINNSVLNTL 143
Db 13 IIGCMAPVSIChELVLLTYSLSVTSQDQIRSANLITENPPDSTV--KLEGSLNAl 70

QY 144 IMLSVIVMTIFLWVLYKRYCYKFIHGNIIMSIIMLLFLFTYIYLGEVLTYKTNVAMDYPT 203
Db 71 VFWVLLIAVTFIIVLFLYYNFTNLKHYMRFSAFFLGTMGGAIFLTSIQHFSIPVSIT 130

QY 204 LLITWVNGAVGMVC1HNGK-PVLYQAYLIM-SALMALVFKYLPENSAVILGAISVY 262
Db 131 CFTLLENFTLGLTSVFAGGIPTVLRCQMYVNGIVVAAWFTK-LPEWTTWFLVALALY 189

QY 263 DLAVLCPKGPLRMLVETQAERNEPIPFYI-----SSAMTVGMAKL 307
Db 190 DLAVLAPGGPLRLLVEASSRDEEL-PAVTFARPTVSSGNORRNROSSRLAVGGGV 248

QY 308 DPSSQGALQLPYDEM----EEDSY-----DSFGEPSPYVEFPPLTGY 347
Db 249 SDGSVELQAVRNHDVNOLGRENSHNMDYNAIAVRDIDNDGDDGNGSRGGLERSPLVGS 308

QY 348 PG-----EE---LEBE-----ERGV 360
Db 309 PSAEHSTSVGRGNMEDRESYMEDEMSPLVLMGWGDNRERAGLESDNYDISNRGI 368

QY 361 KLGQDFIFYSYLVGKAATGSGDWNTTLACFVAILIGLICLTLAYKKAALPALPISI 420
Db 369 KLGQDFIFYSYLVGGRAMY--DLMTVYACI LIISSGLGCTLLSVNRAALPALPISI 425

QY 421 TFGLIFYFSTDNIVRPFM 438
Db 426 MLGVVFFLTRLMEPFV 443

Search completed: March 18, 2000, 14:11:52
Job time: 489 sec

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Sperm membrane protein spe-4 - *Caenorhabditis elegans*
N;Alternate names: probable integral membrane protein
C;Species: *Caenorhabditis elegans*
C;Date: 10-Jun-1993 #sequence_revision 18-Nov-1994 #text_change 09-Sep-1997
C;Accession: A43459; S24632; S24633
R;I;Hernault, S.W.; Andoueno, P.M.
J. Cell Biol. 119, 55-68, 1992

Run on:	March 18, 2000, 16:08:11 ;	Search time 32.57 Seconds					
(without alignments) 410.791 Million cell updates/sec							
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Perfect score:	2356						
Sequence:	1 MLTFMADSDEEEVCDERTSL.....STDNLVRFMDTLASHOLYI 448						
Scoring table:	BLOSUM62						
Searched:	82229 seqs, 29864866 residues						
Database :	SwissProt_38;*						
Word size :	0						
Number of hits that pass the threshold :	82229						
<p>Prd. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.</p> <p>SUMMARIES</p>							
Result No.	Score	Query	Match	Length	DB	ID	Description
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1	2336	100.0	448	1	PSN2_HUMAN	P4910 homo sapien	P33782 antechinus
2	2232	95.5	448	1	PSN2_MOUSE	Q61144 mus musculus	Q34289 dasyurus australis
3	2227	95.3	448	1	PSN2_RAT	Q88777 rattus norvegicus	Q35810 smilichthys
4	2218	94.9	445	1	PSN2_MICM0	P7901 microcebus	P55781 galus morinus
5	1624.5	69.5	449	1	PSN2_XENLA	O12977 xenopus laevis	P13974 escherichia coli
6	1468	62.8	467	1	PSN1_MOUSE	P49169 mus musculus	Q34321 dasyurus haematus
7	1457	62.8	467	1	PSN1_HUMAN	P49168 homo sapiens	
8	1459.5	62.5	468	1	PSN1_RAT	P97387 rattus norvegicus	
9	1449.5	62.1	433	1	PSN1_XENLA	O12976 xenopus laevis	
10	1147	61.9	467	1	PSN1_MICM0	P79002 microcebus	
11	1150	49.2	541	1	PSN1_DROME	Q02092 drosophila	
12	1043	44.6	836	1	YL4_KCAEL	Q20766 caenorhabditis elegans	
13	983	42.1	461	1	SE12_CAEEL	P52266 caenorhabditis elegans	
14	524.5	22.5	358	1	HOP1_CAEEL	Q02100 caenorhabditis elegans	
15	513	22.0	453	1	PSNH_ARATH	Q64638 arabidopsis thaliana	
16	274	11.7	465	1	SPE4_CAEEL	Q01088 caenorhabditis elegans	
17	119	5.1	531	1	YDFG_SCHPO	Q14387 schizosaccharomyces pombe	
18	110	4.7	2016	1	C1NS_HUMAN	Q14544 homo sapiens	
19	109	4.7	1840	1	CIN4_RAT	P15310 rattus norvegicus	
20	108	4.6	381	1	CYB_DASMA	Q34302 dasycercus assatus	
21	104	4.5	1476	1	CFTR_MOUSE	P26361 mus musculus	
22	103	4.4	1450	1	CFTR_RABIT	Q00554 oryctoaguus pilosus	
23	103	4.4	382	1	CYB_DIDMA	P41903 didelphis marsupialis	
24	103	4.4	380	1	CYB_MICL0	P56731 microtus longicaudus	
25	102.5	4.4	893	1	BOSS_DROVI	Q24138 drosophila melanogaster	
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27	102	4.4	381	1	CYB_DASMA	Q03322 dasycercus assatus	
28	102	4.4	381	1	CYB_PSENI	Q35533 pseudantechinus macrourus	
29	101.5	4.3	381	1	CYB_NINNY	Q35196 ningalui yvo	
30	101.5	4.3	381	1	CYB_PARAP	Q35337 parantechinus apicalis	
31	101	4.3	1951	1	CYB_PLAMS	Q3533 planigale macrotis	
32	100.5	4.3	381	1	CYB_PSEMD	Q35333 pseudantechinus macrourus	
33	100.5	4.3	381	1	CYB_DASGE	Q02004 dasycercus geoffroyi	
34	100	4.3	381	1	CIN3_RAT	P15389 rattus norvegicus	
35	99	4.2	381	1	CINS_RAT	Q63334 antechinus stuartii	
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37	99	4.2	381	1	CYB_PHATA	Q4325 homo sapiens	
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ALIGNMENTS

RESULT	1	PSN2_HUMAN	STANDARD;	PRT;	448 AA.
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AC	P49810;				
DT	01-OCT-1996	(Rel. 34, Created)			
DT	01-OCT-1996	(Rel. 34, Last sequence update)			
DT	15-JUL-1999	(Rel. 38, Last annotation update)			
DE	PRESERILIN 2 (PS-2) (STM-2)	(B5-1) (AD3LP)	(AD5).		
GN	PSEN2 OR PSNL2 OR AD4 OR PS2 OR STM2.				
OS	Homo sapiens (Human).				
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
RN	SEQUENCE FROM N.A., AND VARIANT FAD ILE-141.				
RP	SEQUENCE FROM N.A., AND VARIANT FAD ILE-141 AND VAL-239.				
RX	LEVY-LAHAD E., WASCO W., POORAKAJ P., ROMANO D.M., OSHIMA J., PETTINGELL W.H., YU C.-E., JONDRO P.D., SCHMIDT S.D., WANG K., CROWLEY A.C., FU Y.-H., GUENETTE S.Y., GALAS D., NEMENS E., WIJSMAN E.M., BIRD T.D., SCHELLENBERG G.D., TANZI R.E.; "Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Science 269:973-977(1995).				
RN	"Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Nature 376:775-778(1995).				
[2]	SEQUENCE FROM N.A., AND VARIANT FAD ILE-141 AND VAL-239.				
RP	SEQUENCE FROM N.A., AND VARIANT FAD ILE-141 AND VAL-239.				
RX	LEVY-LAHAD E., WASCO W., POORAKAJ P., ROMANO D.M., OSHIMA J., PETTINGELL W.H., YU C.-E., JONDRO P.D., SCHMIDT S.D., WANG K., CROWLEY A.C., FU Y.-H., GUENETTE S.Y., GALAS D., NEMENS E., WIJSMAN E.M., BIRD T.D., SCHELLENBERG G.D., TANZI R.E.; "Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Science 269:973-977(1995).				
RX	"Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Nature 376:775-778(1995).				
RN	SEQUENCE FROM N.A.				
RP	SEQUENCE FROM N.A.				
RX	LEVY-LAHAD E., WASCO W., POORAKAJ P., ROMANO D.M., OSHIMA J., PETTINGELL W.H., YU C.-E., JONDRO P.D., SCHMIDT S.D., WANG K., CROWLEY A.C., FU Y.-H., GUENETTE S.Y., GALAS D., NEMENS E., WIJSMAN E.M., BIRD T.D., SCHELLENBERG G.D., TANZI R.E.; "Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Science 269:973-977(1995).				
RN	"Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Nature 376:775-778(1995).				
[3]	SEQUENCE FROM N.A.				
RP	SEQUENCE FROM N.A.				
RX	LEVY-LAHAD E., WASCO W., POORAKAJ P., ROMANO D.M., OSHIMA J., PETTINGELL W.H., YU C.-E., JONDRO P.D., SCHMIDT S.D., WANG K., CROWLEY A.C., FU Y.-H., GUENETTE S.Y., GALAS D., NEMENS E., WIJSMAN E.M., BIRD T.D., SCHELLENBERG G.D., TANZI R.E.; "Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Science 269:973-977(1995).				
RN	"Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Nature 376:775-778(1995).				
[4]	SEQUENCE FROM N.A.				
RP	SEQUENCE FROM N.A.				
RX	LEVY-LAHAD E., WASCO W., POORAKAJ P., ROMANO D.M., OSHIMA J., PETTINGELL W.H., YU C.-E., JONDRO P.D., SCHMIDT S.D., WANG K., CROWLEY A.C., FU Y.-H., GUENETTE S.Y., GALAS D., NEMENS E., WIJSMAN E.M., BIRD T.D., SCHELLENBERG G.D., TANZI R.E.; "Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Science 269:973-977(1995).				
RN	"Candidate gene for the Alzheimer's disease type 1 familial Alzheimer's disease locus," Nature 376:775-778(1995).				
[5]	SEQUENCE FROM N.A.				
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[6]	VARIANT AD HIS-62.				
RP	VARIANT AD HIS-62.				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
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RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
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RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
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RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M., WEHNERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., CRUTS M., VAN BROEKHOVEN C.; "Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease." Hum. Mutat. 11:183-190(1998).				
RX	CRUTS M., VAN DUIJN C.M., BACKHOVEN H., VAN DEN BROEK M				

Query Match 95.5%; Score 2232; DB 1; Length 448;
 Best Local Similarity 95.5%; Pred. No. 7.3e-14;
 Matches 428; Conservative 6; Mismatches 14; Indels 0; Gaps 0;

Qy 1 M[LF]MADSEEVCDERTSLMSAESPTPSCOBGRQSPEDGNTAQWRSQE[N]EDG[B]DP 60
 Db 1 M[LA]FMADSEEVCDERTSLMSAESPTPSCOBGRQSPEDGNTAQWRTQEESED[C]EDP 60

Qy 61 DRYACSGVGPGRGLEBLTLYGAKHIVMLFVPTLCMIVYYATKS[V]REYTEKNSQLI 120
 Ft 61 DRYACSA[P]GRGLEBLTLYGAKHIVMLFVPTLCMIVYYATKS[V]REYTEKNSQLI 120

Qy 121 YTPFTEDTPSVCQRLLNSVLNTLIMISVIVNTI[F]LVLYKRYC[K]PIHGWLIMSSMLI 180
 Db 121 YTPFTEDTPSVCQRLLNSVLNTLIMISVIVNTI[F]LVLYKRYC[K]PIHGWLIMSSMLI 180

Qy 181 E[L]FTYYLGEVLT[K]TYNAMD[P]TLLT[Q]WNFSAVGMYC[I]HWKGPLVQ[AY]LIMISALMA 240
 Db 181 E[L]FTYYLGEVLT[K]TYNAMD[P]TLLT[Q]WNFSAVGMYC[I]HWKGPLVQ[AY]LIMISALMA 240

Qy 241 LVFI[K]LP[E]WSAWVILGAISYDLYAVLCPKGPLMLVETAOBRNE[P]FPA[L]YSSAMWW 300
 Db 241 LVFI[K]LP[E]WSAWVILGAISYDLYAVLCPKGPLMLVETAOBRNE[P]FPA[L]YSSAMWW 300

Qy 301 TVGMAKLPSSQGALQLPYDPEMEDSYDSFGEPSYDPEVFEPGLTGYPGEELEEBERGV 360
 Db 301 TVGMAKLPSSQGALQLPYDPEMEDSYDSFGEPSYDPEVFEPGLTGYPGEELEEBERGV 360

Qy 361 KLG[GDF]IYSVLGVKAATGGDWNTLACFAILGLCLL[L]A[V]KALPALPISI 420
 Db 361 KLG[GDF]IYSVLGVKAATGGDWNTLACFAILGLCLL[L]A[V]KALPALPISI 420

Qy 421 TFG[LY]FSTDNLVRPMDTLASHOLYT 448
 Db 421 TFG[LY]FSTDNLVRPMDTLASHOLYT 448

RESULT 3
 PSN2_RAT ID PBN2_RAT STANDARD; PRT; 448 AA.
 AC 088777; 035546; 00894 /.
 DT 15-JUL-1999 (Rel. 38, Created)
 DT 15-JUL-1999 (Rel. 38, Last sequence update)
 DT 15-JUL-1999 (Rel. 38, Last annotation update)
 DE PRESENILIN 2 (PS-2)
 GN PSEN2 OR PSEN2 OR PS2.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 RN [1]

SEQUENCE FROM N.A.
 STRAIN=WTSTR; TISSUE=BRAIN;
 EMBL: U57325; AAC53311.1; - .
 EMBL: AF03835; AAB9260.1; - .
 DR [14911]; AAC52935.1; - .
 MGD; MGII:103284; PSEN2.
 PFM; PF01080; Presenilin_1.
 KW Transmembrane; Alternative initiation
 FT CHAIN 1 448 PRESENILIN 2.
 FT CHAIN 298 448 PRESENILIN 2-SHORT.
 FT TRANSMEM 88 106 POTENTIAL.
 FT TRANSMEM 142 160 POTENTIAL.
 FT TRANSMEM 167 188 POTENTIAL.
 FT TRANSMEM 203 219 POTENTIAL.
 FT TRANSMEM 230 246 POTENTIAL.
 FT TRANSMEM 253 269 POTENTIAL.
 FT TRANSMEM 288 305 POTENTIAL.
 FT TRANSMEM 387 406 POTENTIAL.
 FT TRANSMEM 413 429 POTENTIAL.
 FT CONFLICT 87 87 R -> H (IN REF. 2).
 FT CONFLICT 226 226 A -> V (IN REF. 2).
 FT CONFLICT 324 324 MISSING (IN REF. 2).
 SQ SEQUENCE 448 AA; 49955 MW; 680ACF19 CRC32; ;

RP SEQUENCE FROM N.A.
 RC STRAIN=WTSTR; TISSUE=BRAIN;
 RA FRENZEL S.; ABDEL A.S.; LUBBERT H.;
 RA Submitted (JUL-1996) to the EMBL/GenBank/DDJB databases.
 RN [2]

RP SEQUENCE FROM N.A.
 RC STRAIN=WTSTR; TISSUE=BRAIN;
 RX MEDLINE: 97473536.
 RA TAKAHASHI H.; MERCENI M.; NAKAZATO Y.; NOGUCHI K.; MURAYAMA M.;
 RA IMAHORI K.; TAKASHIMA A.;
 RT "Cloning of cDNA and expression of the gene encoding rat presenilin-2."
 RT presenilin-2.
 RL Gene 197:383-387(1997).
 RN [3]

RP SEQUENCE FROM N.A.
 RC STRAIN=WTSTR; TISSUE=BRAIN;
 RX MEDLINE: 98207716.
 RA TANAKA H.; TABIRA T.;
 RT "Cloning of the cDNA encoding rat presenilin-2.";
 RL Biochim. Biophys. Acta 1396:259-262(1998).

|- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE. MAY
 CC SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
 CC FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS (BY SIMILARITY).
 CC -|- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.

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CC DR EMBL; X99267; CAA67663.1;
 DR EMBL; DB3700; BAA2832..1;
 DR EMBL; AB004454; BAA20406.1; - .
 DR PFAM; PF01080; Presenilin_1.
 DR Transmembrane.
 KW Transmembrane.
 FT TRANSMEM 88 106 POTENTIAL.
 FT TRANSMEM 142 160 POTENTIAL.
 FT TRANSMEM 167 188 POTENTIAL.
 FT TRANSMEM 203 219 POTENTIAL.
 FT TRANSMEM 230 246 POTENTIAL.
 FT TRANSMEM 253 269 POTENTIAL.
 FT TRANSMEM 288 305 POTENTIAL.
 FT TRANSMEM 387 406 POTENTIAL.
 FT CONFLICT 7 7 S -> T (IN REF. 1).
 FT CONFLICT 86 87 KH -> ND (IN REF. 3).
 SQ SEQUENCE 448 AA; 50051 MW; 680ACF19 CRC32; ;

Query Match 95.3%; Score 2227; DB 1; Length 448;
 Best Local Similarity 95.3%; Pred. No. 1.6e-14;
 Matches 427; Conservative 6; Mismatches 15; Indels 0; Gaps 0;

Qy 1 MLTFMADSEEVCDERTSLMSAESPTPSCOBGRQSPEDGNTAQWRSQE[N]EDG[B]DP 60
 Db 1 MLAFMADSEEVCDERTSLMSAESPTPSCOBGRQSPEDGNTAQWRTQEESED[C]EDP 60

Qy 61 DRYACSGVGPGRGLEBLTLYGAKHIVMLFVPTLCMIVYYATKS[V]REYTEKNSQLI 120
 Ft 61 DRYACSA[P]GRGLEBLTLYGAKHIVMLFVPTLCMIVYYATKS[V]REYTEKNSQLI 120

Qy 121 YTPFTEDTPSVCQRLLNSVLNTLIMISVIVNTI[F]LVLYKRYC[K]PIHGWLIMSSMLI 180
 Db 121 YTPFTEDTPSVCQRLLNSVLNTLIMISVIVNTI[F]LVLYKRYC[K]PIHGWLIMSSMLI 180

Qy 181 E[L]FTYYLGEVLT[K]TYNAMD[P]TLLT[Q]WNFSAVGMYC[I]HWKGPLVQ[AY]LIMISALMA 240
 Db 181 E[L]FTYYLGEVLT[K]TYNAMD[P]TLLT[Q]WNFSAVGMYC[I]HWKGPLVQ[AY]LIMISALMA 240

Qy 241 LVFI[K]LP[E]WSAWVILGAISYDLYAVLCPKGPLMLVETAOBRNE[P]FPA[L]YSSAMWW 300
 Db 241 LVFI[K]LP[E]WSAWVILGAISYDLYAVLCPKGPLMLVETAOBRNE[P]FPA[L]YSSAMWW 300

Qy 301 TVGMAKLPSSQGALQLPYDPEMEDSYDSFGEPSYDPEVFEPGLTGYPGEELEEBERGV 360
 Db 301 TVGMAKLPSSQGALQLPYDPEMEDSYDSFGEPSYDPEVFEPGLTGYPGEELEEBERGV 360

Qy 361 KLG[GDF]IYSVLGVKAATGGDWNTLACFAILGLCLL[L]A[V]KALPALPISI 420
 Db 361 KLG[GDF]IYSVLGVKAATGGDWNTLACFAILGLCLL[L]A[V]KALPALPISI 420

Qy 421 TFG[LY]FSTDNLVRPMDTLASHOLYT 448
 Db 421 TFG[LY]FSTDNLVRPMDTLASHOLYT 448

RESULT 3
 PSN2_RAT ID PBN2_RAT STANDARD; PRT; 448 AA.
 AC 088777; 035546; 00894 /.
 DT 15-JUL-1999 (Rel. 38, Created)
 DT 15-JUL-1999 (Rel. 38, Last sequence update)
 DT 15-JUL-1999 (Rel. 38, Last annotation update)
 DE PRESENILIN 2 (PS-2)
 GN PSEN2 OR PSEN2 OR PS2.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 RN [1]

Query Match 95.3%; Score 2227; DB 1; Length 448;
 Best Local Similarity 95.3%; Pred. No. 1.6e-14;
 Matches 427; Conservative 6; Mismatches 15; Indels 0; Gaps 0;

Qy 1 MLTFMADSEEVCDERTSLMSAESPTPSCOBGRQSPEDGNTAQWRSQE[N]EDG[B]DP 60
 Db 1 MLTFMADSEEVCDERTSLMSAESPTPSCOBGRQSPEDGNTAQWRTQEESED[C]EDP 60

Qy 61 DRVCGSGVGPGRGLEELTLYGAKHIVMLFVPTLCMIVYYATKS[V]REYTEKNSQLI 120
 Db 61 DHVACSGVGPGRGLEELTLYGAKHIVMLFVPTLCMIVYYATKS[V]REYTEKNSQLI 120

Qy 121 YTPFTEDTPSVCQRLLNSVLNTLIMISVIVNTI[F]LVLYKRYC[K]PIHGWLIMSSMLI 180
 Db 121 YTPFTEDTPSVCQRLLNSVLNTLIMISVIVNTI[F]LVLYKRYC[K]PIHGWLIMSSMLI 180

Qy 181 E[L]FTYYLGEVLT[K]TYNAMD[P]TLLT[Q]WNFSAVGMYC[I]HWKGPLVQ[AY]LIMISALMA 240
 Db 181 E[L]FTYYLGEVLT[K]TYNAMD[P]TLLT[Q]WNFSAVGMYC[I]HWKGPLVQ[AY]LIMISALMA 240

ROYTA M., LILLIUS L., EEROLA A., ST GEORGE-HYSLOP P.H., FREY H., LAUNFELD L.;
"The Glu18Gly mutation of the presenilin-1 gene does not necessarily cause Alzheimer's disease";
Ann. Neurol. 44:965-967(1998). [15]

VARIANT GLY-318
MEDLINE: 99066775.
RA ALDUDO J., BULLIDO M.J., FRANK A., VAIDIVIESO F.;
"Missense mutation E318G of the presenilin-1 gene appears to be a nonpathogenic polymorphism";
Ann. Neurol. 44:985-986(1998). [16]

VARIANTS AD VAL-79; CYS-115; VAL-231, AND VARIANT GLY-318.
MEDLINE: 98046005.
RA CRUTS M., VAN DUTIN C.M., BACKHOVEN H., VAN DEN BROECK M., WEINERT A., SERNEELS S., SHERRINGTON R., HUTTON M., HARDY J., ST GEORGE-HYSLOP P.H., HOFMAN A., VAN BROEKHOVEN C.;
"Estimation of the genetic contribution of presenilin-1 and -2 mutations in a population-based study of presenile Alzheimer disease";
Hum. Mol. Genet. 7:43-51(1998). [17]

VARIANTS AD D-120; R-163; V-209; V-260; L-264; Y-410 AND P-426.
MEDLINE: 98180720.
RA POORKAJ P., SHARMA V., ANDERSON L., NEMENS E., ALONSO M.E., ORR H., WHITE J., HESTON L., BIRD T.D., SCHELENBERG G.D.;
"Missense mutations in the chromosome 14 familial Alzheimer's disease presenilin 1 gene";
Hum. Mutat. 11:216-221(1998). [18]

VARIANT AD GLU-378.
MEDLINE: 99211215.
RA BESANCON R., LORENZI A., CRUTS M., RADAWIEC S., STURTZ F., BROUSSOLE E., CHAZOT G., VAN BROEKHOVEN C., CHAMBA G., VANDENBERGHE A.;
"Missense mutation in exon 11 (codon 378) of the presenillin-1 gene in a French family with early-onset Alzheimer's disease and transmission by mismatch enhanced allele specific amplification";
Hum. Mutat. 11:481-481(1998). [19]

VARIANTS AD LEU-169 AND GLN-436.
MEDLINE: 99047368.
RA TADDEI R., KWOK J.B., KRIL J.J., HALLDAY G.M., CREASEY H., HALLUPP M., FISHER C., BROOKS W.S., CHUNG C., ANDREWS C., MASTERS C.L., SCHOFIELD P.R., MARTINS R.N.;
"Two novel presenilin-1 mutations (Ser169Leu and Pro436Gln) associated with very early onset Alzheimer's disease";
NeuroReport 9:3335-3339(1998). [20]

VARIANT AD PRO-169.
MEDLINE: 99148656.
RA EZQUERA M., CARNERO C., BLESA R., GELPI J.L., BALLESTA F., OLIVA R.;
"A presenilin 1 mutation (Ser169Pro) associated with early-onset AD and myoclonic seizures";
Neurology 52:566-570(1999).

RA	QY	188	LGEVLKTYNAMDYPTLLTVNFGAVGMVCIHWKGPLVQAYLIMITSALMVFKYL 247
RA	Db	182	LGEVFKTYNADVDTIYVMTSHWKGPLRQAYLIMITSALMVFKYL 241
RT	QY	248	PENSAWTLGAIISVYDLVAVLCPKGPFLRMVLYTATKSYFVYGMVTSIHWKGPLRQAYLIMITSALMVFKYL 307
RT	Db	242	PENAWLILAVLSVIDLVAVLCPKGPFLRMVLYTATKSYFVYGMVTSIHWKGPLRQAYLIMITSALMVFKYL 301
RP	QY	308	DPSQQAL-QLPYDP-E-MEEDSYDFGE--PSYPYEFEPPLTGYPG-----350
RP	Db	302	DPEAQRRVSKNSKNAESTEREQDVAENDDGGESEWEAQDSDHGPHRSTPESRAA 361
RP	QY	350	EEL-----EEPEERVKLGQDFIFIVSVLVKGAAATSGDWNTLACFVALIGCLT 402
RP	Db	362	QEISSSLAGEPEEYKLGQDFIFIVSVLVKGAAATSGDWNTLACFVALIGCLT 421
RX	QY	403	LLILAVFKKALPALPISITFGLLIFYFSTDLYRPMDTLASHOLYI 448
RA	QY	422	LLILIAFKKALPALPISITFGLLIFYFATDLYQPFMDOLAFHQFYI 467
<hr/>			
RESULT 8			
PSNL_RAT	ID	P91871_RAT	STANDARD;
PSNL_RAT	AC	P91871; P97529;	PRT;
PSNL_RAT	DT	15-JUL-1999 (Rel. 38, Created)	
PSNL_RAT	DT	15-JUL-1999 (Rel. 38, Last sequence update)	
PSNL_RAT	DT	15-JUL-1999 (Rel. 38, Last annotation update)	
PSNL_RAT	DE	PRESENTIN 1 (PS-1) (S182 PROTEIN).	
PSNL_RAT	GN	Rattus norvegicus (Rat).	
PSNL_RAT	OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.	
PSNL_RAT	RN	[1]	
PSNL_RAT	RP	SEQUENCE FROM N.A.	
PSNL_RAT	RC	STRAININWISTAR; TISSUE-BRAIN;	
PSNL_RAT	RX	MEDLINE: 97199371.	
PSNL_RAT	RA	TANIGUCHI T., HASHIMOTO T., TANIGUCHI R., SHIMADA K., KAWAMATA T., YASUDA M., NAKAI M., TERASHIMA A., KOIZUMI T., MAEDA K., TANAKA C.; Cloning of the cDNA encoding rat presenilin-1.";	
PSNL_RAT	RA	Gene 186:73-75(1997).	
PSNL_RAT	RL	[2]	
PSNL_RAT	RN	SEQUENCE FROM N.A.	
PSNL_RAT	RC	STRAININWISTAR; TISSUE-BRAIN;	
PSNL_RAT	RX	MEDLINE: 96255262.	
PSNL_RAT	RA	TAKAHASHI H., MURAYAMA M., TAKASHIMA A., MERCKEN M., NAKAZATO Y., NOGUCHI K., IMAHORI K.; Molecular cloning and expression of the rat homologue of presenilin-1.";	
PSNL_RAT	RT	Neurosci. Lett. 206:113-116(1996),	
PSNL_RAT	CC	-1- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE.	
PSNL_RAT	CC	CC the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement. (See http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).	
PSNL_RAT	CC	-1- MAY FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS.	
PSNL_RAT	CC	-1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.	
PSNL_RAT	CC	-1- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.	
PSNL_RAT	CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - CC the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement. (See http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).	
PSNL_RAT	DR	EMBL: D82578; BAA11575_1;	
PSNL_RAT	DR	DR EMBL: D82363; BAA11564_1;	
PSNL_RAT	DR	PPBM: PF01080; Presenilin 1.	
PSNL_RAT	KW	Transmembrane; Glycoprotein.	
PSNL_RAT	FT	Transmem 8 103	POTENTIAL.
PSNL_RAT	FT	Transmem 13 3 153	POTENTIAL.
PSNL_RAT	FT	Transmem 16 1 181	POTENTIAL.
PSNL_RAT	FT	Transmem 19 5 215	POTENTIAL.
PSNL_RAT	FT	Transmem 22 1 241	POTENTIAL.

Query Match Score 1467; DB 1; Length 467;
Best Local Similarity 62.8%; Pred. No. 3-5e-94;
Matches 306; Conservative 39; Mismatches 79; Indels 42; Gaps 8;

QY 24 ESPTPRSCQEGRQGPEDGENTAQWRSQNEEDGEEDPDRYVCSGVP----GRPPG----75
Db 3 ELPAPLSYFQNAQMSEDNHLSNTVRSDNRQRERHDDR-SLGHPEPLSNRQNSRQ 61

QY 75 -----LEELLTLYGAKHYTMFLFVPTLCKMIVVATKSYFTEKNGQLIYTPFTED 127

Db 62 YVEQDEEEDEELTLKYGAKHYTMFLFVPTLCKMIVVATKSYFTEKNGQLIYTPFTED 121

QY 128 TPSVGQRLNLSVNTLMISVVMFTFLVWLYKRYCFTHGLWIMSLMLFLTYII 187

Db 122 TETVGQRLAHSLINAAMISVIVMFTLWVLYKRYCVTHAWLWLISSLFFFSFIY 181

PSNL_MICOMU	STANDARD;	PRT;	467 AA.
ID P78802			
DT 15-JUL-1999 (Rel. 38, Created)			
DT 15-JUL-1999 (Rel. 38, Last sequence update)			
DT 15-JUL-1999 (Rel. 38, Last annotation update)			
DE PRESENILIN 1 (PSL-1).PSI.			
OS Microcebus murinus ('lesser mouse lemur').			
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;			
OC Eutheria; Primates; Strepsirrhini; Cheirogaleidae; Microcebus.			
RN [1]			
RP SEQUENCE FROM N.A.			
TISSUE="BRAIN";			
RX MEDLINE: 97079199.			
RA CALENDA A., MESTRE-FRANCES N., CZECH C., PRADIER L., BONS N.,			
RA BELLIS M.;			
RT "Molecular cloning, sequencing, and brain expression of the			
presenilin 1 gene in <i>Microcebus murinus</i> ".			
RT Biochim. Biophys. Res. Commun. 248:430-439 (1996).			
CC -!- FUNCTION: MAY PLAY A ROLE IN INTRACELLULAR SIGNALING AND GENE EXPRESSION OR IN LINKING CHROMATIN TO THE NUCLEAR MEMBRANE.			
CC MAY FUNCTION IN THE CYTOPLASMIC PARTITIONING OF PROTEINS.			
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.			
CC -!- ALTERNATIVE PRODUCTS: TWO ISOFORMS; I-467 (SHOWN HERE) AND I-463 ARE PRODUCED BY ALTERNATIVE SPLICING.			
CC -!- TISSUE SPECIFICITY: FOUND PREDOMINANTLY IN NEURONS OF THE DIFFERENT CORTICAL LAYERS AND HIPPOCAMPUS BUT ALSO IN SUBCORTICAL STRUCTURES.			
CC -!- SIMILARITY: BELONGS TO THE PRESENILIN FAMILY.			
CC -----			
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CC -----			
EMBL: 271333; CAA95930.1; -;			
PFAM: PF01080; Presenilin_1.			
KW Transmembrane; Alternative splicing; Glycoprotein.			
FT TRANSMEM 83	103		
FT TRANSMEM 133	153	POTENTIAL.	
FT TRANSMEM 161	181	POTENTIAL.	
FT TRANSMEM 191	211	POTENTIAL.	
FT TRANSMEM 221	241	POTENTIAL.	
FT TRANSMEM 244	264	POTENTIAL.	
FT TRANSMEM 281	301	POTENTIAL.	
FT TRANSMEM 408	428	POTENTIAL.	
FT TRANSMEM 433	453	POTENTIAL.	
FT CARBOHYD 279	279	POTENTIAL.	
FT VARSPLC 405	405	POTENTIAL.	
FT VARSPLC 26	29	MISSING (IN ISOFORM I-463).	
SQ SEQUENCE 467 AA;	52384 MW;	A841AOB7 CRC32;	
CC -----			
Query Match Score 1447; DB 1; Length 467;			
Best Local Similarity 64.4%; Pred. No. 8.4e-93; 7;			
Matches 300; Conservative 41; Mismatches 83; Indels 42; Gaps 7;			
Qy 24 ESTTPRSQCEGRGPEDGENTAGWRSOENEDGEEDPDRYCSGVP-----GRPPG---- 75			
Db 3 ELPAPLSYFQNMQNSDHLNSNTVRSDQDNREQDHGRRRL-GNPEPLSNGPQGNSGP 61			
Qy 75 -----LEELLTLYGAKHIVMLFYPVTLCKMIVVATIKSYFTEKNGOLIYTPPTED 127			
Db 62 VVERDEEEDEELTLYGAKHIVMLFYPVTLCKMIVVATIKSYFTEKNGOLIYTPPTED 121			
Qy 128 TPSVGQLLNNTVNTLMISIVVMTFLVVLKYRCVKFIHQWLNSMLFLFYIY 187			
Db 122 TEPVGQRLHSVNLAIMSVIVMTFLVVLKYRCVKIHWLISLLFFSFYI 181			
KW Transmembrane; Glycoprotein; Alternative splicing.			
FT TRANSMEM 107 127 POTENTIAL.			

SEL-12. *Caenorhabditis elegans*.
 Nematoda: Secernentea; Rhabditida;
 Eukaryota; Metazoa; Rhabditina; Rhabditidae; Peioderinae; Caenorhabditis.
 [1]
 SEQUENCE FROM N.A.
 STRAIN-BRISPOL N2.
 MEDLINE: 9602531.
 LEVITAN D., GREENWALD I.;
 "Facilitation of lin-12-mediated signalling by sel-12,
Caenorhabditis elegans S182 Alzheimer's disease gene.",
Nature 377:351-354 (1995).

REVISIONS TO 84-85.

LEVITAN D.; Submitted (JAN-1996) to the EMBL/GenBank/DDBJ databases.

-|- FUNCTION: MAY FACILITATE LIN-12 MEDIATED RECEPTION OF INTERCELLULAR SIGNALS. IT MIGHT BE DIRECTLY INVOLVED IN LIN-12 MEDIATED RECEPTION, FUNCTIONING AS A CO-RECEPTOR OR AS A DOWNSTREAM EFFECTOR THAT IS ACTIVATED UPON LIN-12 ACTIVATION.

-|- ALTERNATIVELY IT MAY BE INVOLVED IN A MORE GENERAL CELLULAR PROCESS SUCH AS RECEPTOR LOCALIZATION OR RECYCLING AND HENCE INFLUENCE LIN-12 ACTIVITY INDIRECTLY.

-|- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.

-|- SIMILARITY: BELONGS TO THE PRENINILIN FAMILY.

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	POTENTIAL.	POTENTIAL.	POTENTIAL.	POTENTIAL.	C->S: ARI31; EGG-LAYING-DEFECTIVE.
TRANSMEM	46	66			
TRANSMEM	102	122			
TRANSMEM	131	151			
TRANSMEM	164	184			
TRANSMEM	190	210			
TRANSMEM	213	233			
TRANSMEM	251	271			
TRANSMEM	385	405			
MUTAGEN	60	60			
SEQUENCE	461	AA;	51817 MW;	ABDD085 CRC32;	

Dy	26	PTRPRSCQEGRQRGPEDGE - NTAQORSOENEEEDGEEDPDRYVCSPGPRPGLEELTLKYG	84
Db	2	PSTRRQEG - -GGADAETHVTVGTNLITNRSQEDEN - - - - - - - - - - - - - - -	46
Dy	85	AKHVIMLEVFPVTLCMIVVATIKSVRFITEKNGO - LIVTPFTEDTPSVQRLLASVLNTL	143
Db	47	ASHVILHFVPUVSCLUMVFTTNTIIFYSNQNRGRHLLYTFPVRETDSYIEFKGLMSLGNAL	106
Dy	144	TMISVTTMFITFLVYKRYCFKIGWLMISLMLFLFTYIYLGEVKRTYNAAMDPT	203
Db	107	WNLCCVVUMLMTLVLLIVPKYKFKLIGHNLIVSSSTLLFETTIVQEVIKSFDVSPSALL	166
Dy	204	ILLTYWNFGAVGMVCHWKGPFLVLLQQAYLIMISALMALVYKYLPEWSAWVILGAISYVD	263
Db	167	VFLGLGNYGVLMCWHKGPMCHWLQQLFLITMSALMALVYKYLPEWTWFLVEVISWVD	226
Dy	264	L-YAVLCPKGPLRMLIVEAQERNEPIFPALIYSSAMWTVTGMAKLDPSOGAQLPYDPEN	323
Db	227	IYAVLTKEGPRLYLVAAERNEPITPAPLTYSSCSEVYPPVTVT	279

Query Match 22.5%; Score 524.5; DB 1; Length 358;
 Best Local Similarity 30.5%; Pred. No. 2.1e-29;
 Matches 117; Conservative 81; Mismatches 136; Indels 49; Gaps 0
 83 YGAKHYIMLFYPVTLCMIVVATKSVREYTEKNGOLITYTPF---TEDTPSYGGORLNLNS 138

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: March 18, 2000, 22:05:40 ; Search time 49.26 Seconds
 (without alignments)
 630.566 Million cell updates/sec

Title: US-08-509-359B-137
 Perfect score: 2336
 Sequence: 1 MLTFMASDSEEEVCDERTSL.....STDNLVRPFMDTLASHOLYI 448

Scoring table: BIOSUM62
 Searched: 225878 seqs, 69334122 residues
 Database : SPTREMBL_12:
 Word size : 0

Number of hits that pass the threshold : 225878

1: sp_archaea:*

2: sp_bacteria:*

3: sp_fungi:*

4: sp_human:*

5: sp_invertebrate:*

6: sp_mammal:*

7: sp_mhc:*

8: sp_organelle:*

9: sp_phage:*

10: sp_plant:*

11: sp_rhodent:*

12: sp_virus:*

13: sp_vertebrate:*

14: sp_unclassified:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,

and is derived by analysis of the total score distribution.

SUMMARIES

Result	No.	Score	Query	Match	Length	DB	ID	Description
-	1	2262.5	96.9	449	6	Q9XT96		Q9xt96 bos taurus
	2	1438.6	61.6	478	6	Q9XT97		Q9xt97 bos taurus
	3	1407.5	60.3	456	13	Q9W6T7		Q9wt7 brachydanio
	4	1355.5	58.0	384	13	Q73869		Q73869 cyprinus ca
	5	576.5	24.7	272	5	Q96340		Q96340 drosophila
	6	440.0	18.8	184	4	Q95465		Q95465 homo sapien
	7	113.5	4.9	406	5	Q19737		Q19737 caenorhabdi
	8		4.8	4578	13	Q42181		Q42181 fugu rubrip
	9	110.5	4.7	320	8	Q34086		Q34086 coecyulus er
	10	110.5	4.6	381	8	Q35425		Q35425 phascolosor
	11	109	4.7	1840	11	P07611		P07611 rattus norv
	12	108.5	4.6	380	8	Q9ZC9		Q9zc9 upupa
	13	107.5	4.6	748	2	Q92577		Q92577 streptomyce
	14	107	4.6	381	8	Q33723		Q33723 antechinus
	15	106.5	4.6	382	8	Q34677		Q34677 didelphis a
	16	106.5	4.6	318	11	Q35294		Q35294 rattus norv
	17	105	4.5	303	11	P97829		P97829 rattus norv
	18	106	4.5	652	5	Q93346		Q93346 caenorhabdi
	19	106	4.5	382	8	Q34279		Q34279 dielephius a
	20	106	4.5	1717	13	Q90519		Q90519 fugu rubrip
	21	106	4.5	444	4	Q9X2N3		Q9x2n3 arthropoete
	22	105	4.5	379	8	Q34428		Q34428 echimys did
	23	104	4.5	382	8	Q35561		Q35561 philander o
	24	104	4.5	379	8	Q36096		Q36096 trinomys pa

ALIGNMENTS

RESULT	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	
ID	Q9XT96																									
AC	Q9XT96;																									
DT	01-NOV-1999	(TREMBLrel.	12,	Created)																						
DT	01-NOV-1999	(TREMBLrel.	12,	Last sequence update)																						
DE	PRESENTILIN	2.																								
OS	Bos taurus (Bovine).																									
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;																									
OC	Butheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovidae;																									
RN	[1]																									
RP	SEQUENCE FROM N.A.																									
RC	TISSUE-BRAIN;																									
RA	SAHARA N., SHIRASAWA T., MORI H.;																									
RT	"Molecular cloning of bovine presenilin 2 gene."																									
RL	"Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.																									
DR	EMBL; AF03937; AAD30241;																									
SO	SEQUENCE 44 AA;																									
	50301 MW;																									
	A3DR878F CRC32;																									

Query	Match	96.9%	Score	2262.5;	DB	6;	Length	449;
Best Local Matches	436;	Conservative	5;	Mismatches	7;	Indels	1;	Gaps 1;
1	MLTFMASDSEEVCDERTSLMSAESPTRPSQEGRGPEDGENTAOWRSQNEEND-SEED	59	Qy					
2	MLTFMASDSEEVCDERTSLMSAESPTRPSQCDGROGLDGESEAQWRSQSEEDHEED	60	Db					
3	PDRYVGSPGRPPGLEELTKYGAHKYMLFVFPVTLCMVVATIKSVRFYTEKNGQL	119	Qy					
4	LFLFTTYLGEVLTKYNAVDYPTLLTLYKAVHVNDFVNGMCIHWKGPLVLYQQAYLIMISALM	239	Db					
5	PDRYVGSPGRPPGLEELTKYGAHKVNLFVFPVTLCMVVATIKSVRFYTEKNGQL	120	Qy					
6	IYPFEDTPSYGQRLNSVNTLIMISIVMTIPLVLYKRYCKFTHGWLIMSSML	179	Qy					
7	IYPFEDTPSYGQRLNSVNTLIMISIVMTIPLVLYKRYCKFTHGWLIMSSML	180	Db					
8	ALVFIXKLPERSAWVYLIGAISYDLYAVLCPKGLMVEAQERNPFPALIYSSAMY	299	Qy					
9	ALVFIXKLPERSAWVYLIGAISYDLYAVLCPKGLMVEAQERNPFPALIYSSAMY	300	Db					
10	WTGMAKLDPSSQGALQLPYSPEMEYDSYDSTFGEPSYVFPFPPLTGYPGEELEEE	359	Qy					

RN [1]	SEQUENCE FROM N.A.	Qy 240 ALVFIKYLPENSAWILGAISYDLYAVLCPKGLPRLMIVTAQERNEPIFPALEYSSAMV 299
RP ARCHER S., HITANO J., DISS J.K., FRASER S.P., DURANGOZ M.B.A.;	Db 1 ALVFIKYLPENSAWILGAISYDLYAVLCPKGLPRLMIVTAQERNEPIFPALEYSSAMV 60	
RA RL NeuroReport 0-0(0).	Qy 300 WTV-----GMAKLDPSS-----QGAIQLQYDPEMEEDSYDSFGEPYPE- 339	
DR EMBL; Y7128; CAA76641.1;	Db 61 YALVNNTVTPOOSQATASSPNSNTTTRATQNSLA--SPEAAASQRTGN-SHPRQ 116	
PFAM; PF01080; Presenilin; 1.	Qy 339 -----WPEPLTGPGE--ELEE----- 355	
FT NON_TER 1	Db 117 NORDGSVLAATEAAGFTQEWANSIERYARROQEVOSTQSGNAQRSEYRTVAPDQN 176	
SEQUENCE 384 AA; 43276 MW; 21A78D17 CRC32;	Qy 355 -----EBERGKYLGLDFIFTSVLYGRAAATGSDGNTTLACFVATLGCLTLLIAFK 410	
Query Match 58.0%; Score 1355; DB 13; Length 384;	Db 177 HDGQPERGKFLGLGFIFTSVLYGRAS--YGDWTIACPVAILGCTTLLIAWR 234	
Best Local Similarity 69.9%; Pred. No. 3.9e-94;	Db 117 KALPALPISITFGFLIEFYSTDNLVRPEMDTLASHOLYI 448	
Matches 270; Conservative 33; Mismatches 55; Indels 28; Gaps 5;	Qy 411 KALPALPISITFGFLIEFYSTDNLVRPEMDTLASHOLYI 448	
Qy 89 IMLFIPVPTLCMVYVATKSVPVFTEKNGQLIYTFPDTPSYQRLINSLVNTLIMS 147	Db 235 KALPALPISITFGFLIEFCATSAYVKEFMDLSAKOFEI 272	
1 IMLFIPVPTLCMVYVATKSVPVFTEKNGQLIYTFPDTPSYQRLINSLVNTLIMS 60	RESULT 6	
Db 181 LCPKGPLRILVETQAERNPFPFALIYSSMFTVQRLNSMLNRAIMTS 60	095465 PRELIMINARY; PRT: 184 AA.	
Qy 148 VIVVNTIELVLYLVKRYCYRFIGWLMSSLFLFIFTYLYGEVLRKTYNAMDYPFLLT 207	ID 095465 ID 095465 PRELIMINARY; PRT: 184 AA.	
61 VIVVNTIELVLYLVKRYCYRVIGLEFFSEFYLGFEKTYNAMDYFLAVI 120	AC 095465 DT 01-MAY-1999 (TREMBLrel. 10, Created)	
Qy 208 VWNFGAVGMYCINWKGPVLQOALIMISALMALVFIKYLPENSAWILGAISYDLYAV 267	DR 01-MAY-1999 (TREMBLrel. 10, Last sequence update)	
Db 121 IWFNSVVGMCINWKGPRLQOALIMISALMALVFIKYLPENSAWILGAISYDLYAV 180	DB 01-MAY-1999 (TREMBLrel. 10, Last annotation update)	
Qy 268 LCPKGPLRILVETQAERNPFPFALIYSSMFTVQRLNSMLNRAIMTS 60	DE MINILIN.	
Db 181 LCPKGPLRILVETQAERNPFPFALIYSSMFTVQRLNSMLNRAIMTS 60	GN OS Homo sapiens (Human)	
Qy 319 -----YDPMEEEDSYDSFG-----EPSYVEPEPLTGPGEDEEBERGVKL 362	OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Primates; Catarrhini; Hominoidea; Homo.	
Db 241 APTAQPEDDGFTPAWVQQHQPMQSTEDSRREIQELPSAPP- PVEDDEERGVKL 298	OC Buteraria; Primates; Catarrhini; Hominoidea; Homo.	
Qy 363 GLGDEFITFSYLSVYGRAATGSDGNTTLACFVATLGCLTLLIAVEKKALPALPISITE 422	RN [1] SEQUENCE FROM N.A.	
Db 299 GLGDEFITFSYLSVYGRAATGSDGNTTLACFVATLGCLTLLIAVEKKALPALPISITE 358	RP GEGG M.E.; PALMER M.S.; POWELL C.S.	
Qy 423 GLIIFYSTDNLYVRPEMDTLASHOLYI 448	RA Human presenilin 1 gene encodes an alternative protein-minilin."	
Db 359 GLVFFYFATDNLYVRPEMDQLAHVQFYI 384	RT Submitted (AUG-1998) to the ENBL/GenBank/DDJB databases.	
RESULT 5	RL EMBL; AJ000005; CAAU7825.1; DR AJ000005; CAAU7825.1; SQ SEQUENCE 184 AA; 21073 MW; 5C6FBAEE CRC32;	
O96340 096340 PRELIMINARY; PRT: 272 AA.	Query Match 18.8%; Score 440; DB 4; Length 184;	
ID 096340 PRELIMINARY; PRT: 272 AA.	Best Local Similarity 56.0%; Pred. No. 6.2e-26; Matches 102; Conservative 15; Mismatches 43; Indels 22; Gaps 4;	
AC 096340; DT 01-MAY-1999 (TREMBLrel. 10, Created)	Qy 24 ESPTPRSCOEGRGPEDGENTAOQWSQENEEDGEEDPDYYCSGYP----GRPPG --- 75	
DT 01-MAY-1999 (TREMBLrel. 10, Last sequence update)	Db 3 ELPAPLSPYFQNAQMSEDDNHLISNTVRSQNDNRERQEHDRR-SLGHEPELSNRPQNSRQ 61	
DE PRESENTILIN (FRGMENT).	Qy 75 -----LEEEETLKYGAKHIMLFPTVLCMIVVATIKSVPRTYERKGQLIYTPTED 127	
OS Drosophila melanogaster (Fruit fly).	Db 62 VVEQDEEDEETLKYGAKHIMLFPTVLCMIVVATIKSVPRTYERKGQLIYTPTED 121	
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephdroioidea; Drosophilidae; Drosophila.	Qy 128 TPSYGQLRLNSVNTLIMISYIVVMTFLFVLYKRYCKYKFIHGWLMSLMLFLTYIY 187	
RP SEQUENCE FROM N.A.	Db 122 TETVGORALHSILNAIMISVIVVMFLVVLYKRYCKYKFIHGWLMSLMLFLTYIY 176	
RC STRAIN=CANTON S;	Qy 188 LG 189	
RX MEDLINE: 98331525.	Db 177 LG 178	
RA MARFANY G., DEL-FAVERO J., VALERO R., DE JONGHE C., WOODROW S., HENDRIKS L., VAN BROECKHOVEN C., GONZALEZ-DUARTE R.;	RESULT 7	
FT NON_TER 1	Q19737 ID Q19737 PRELIMINARY; PRT: 406 AA.	
SEQUENCE 272 AA; 29456 MW; 606B9A5C CRC32;	AC Q19737 DT 01-NOV-1996 (TREMBLrel. 01, Created)	
Query Match 24.7%; Score 576.5; DB 5; Length 272;	DR F22E10.5 PROTEIN.	
Best Local Similarity 47.8%; Pred. No. 5.7e-36;	SQ DE F22E10.5.	
Matches 133; Conservative 29; Mismatches 41; Indels 75; Gaps 8;	GN Caenorhabditis elegans.	

Qy	97	LCKMIVVATIKSVRFYTERNGQOLITYPFTDTPSYGQRLLNSVLNTLIMISIVVMTIFL	156
:	:	:	
Db	7	LCLLTQIVT-----GLLFLAMHYTADT-----	35
Qy	157	VVLYKRYCKFINGWLI----MSSMLLELFETYIYIGEVLKTYNNAMDYPILLTNWFG	212
: :	: :	: :	
Db	35	-----AHTCENVOQGWLRNLHANGASSMFVICIYIHLGRGF-YYGSYLNKET-----WNTG	84
Qy	213	AVGMVCITHRKGPVLQOAYLIMISALMAYEIKYDPEWSAWVILGMNSVYDLYAVLPKG	272
: :	: :	: :	
Db	85	-----VILLLTMAFATPGVYLPWGMSEFWGTVITFSAPFYIG	125
Qy	273	PLRMLVETAQ---ERNEP-----1FPALIYSAMYWTVGMPAKLDPSSQGA Q---	317
: :	: :	: :	
Db	126	-OTLVENAWGGFSVDNPNTLTFTRFLAHFLLPFMIAQLTIVHLTFLETGSNNPLGQSNC	183
Qy	317	-LRYDPMEEDSYDSFG-----EPSY---PEFVEP---PLTGYPGEBELDEEE	357
: :	: :	: :	
Db	184	DK1SFHPYPSLKDVLGVFTTLILLTALFSPTLGDPEFSPANLVTP-----	235
Qy	358	RGYKLGLGDEFIYSVLGKAATGSGDWNTLACAFAILGLCLTLLLAYFKKALPALP	417
: :	: :	: :	
Db	235	-HVKPDWSEFFETATRSPIKLG-----VLAIRKESVY-LSLAPLLKNSKKRAMPSRP	287
Qy	418	IISTFGFLIFYFSTDNLVRFEMDTFLASH	444
: :	: :	: :	
Db	288	LSQT---LFWFLVNF--ILKWLGSH	309
RESULT 10			
Q35425		PRELIMINARY;	PRT; 381 AA.
ID	Q35425;		
AC	Q54425;		
DT	01-NOV-1996	(TREMBLE 01, Created)	
DT	01-MAY-1999	(TREMBLE 10, Last sequence update)	
DT	01-MAY-1999	(TREMBLE 10, Last annotation update)	
DE	CYTOCROME C B.		
OS	Phascolosorex dorsalis.		
OG	Mitochondrion.		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;		
RN	[1]		
RP	SEQUENCE FROM N.A.		
RX	MEDLINE; 93096825.		
RA	KRAJEWSKI C, DRISKELL A.C., BAVERSTOCK P.R., BRAUN M.J.;		
RT	"Phylogenetic relationships of the thyaciine (Mammalia: Thylaciniidae) among dasyuroid marsupials: evidence from cytochrome b DNA sequences."		
RT	Proc. R. Soc. Lond., B, Biol. Sci. 250:19-27(1992).		
RL	-I - CATALYTIC ACTIVITY: QH(2) + 2 FERRICYTOCHROME C = Q + 2		
CC	FERRICYTOCHROME C.		
CC	- I - COFACTOR: TWO HEME GROUPS (B562 AND B566) WHICH ARE NOT COVALENTLY BOUND TO THE PROTEIN (BY SIMILARITY).		
CC	EMBL: M99462; AAB61692.1; -.		
DR	PF00033; cycochrome_b_N; 1.		
DR	Mitochondrion; Electron transport; Respiratory chain; Transmembrane; Hem.		
KW	Sequence		
SQ	381 AA; 42813 MW;	C6A2674F CRC32;	
Query Match Score 4.7%; Best Local Similarity 21.1%; Matches 81; Conservative 54; Mismatches 115; Indels 133; Gaps 1			
Qy	97	LCMIVVATIKSVRFYTERNGQOLITYPFTDTPSYGQRLLNSVLNTLIMISIVVMTIFL	156
: :	: :	: :	
Db	39	MCLLQIQT-----GLFLAMHYTSD-----	64
Qy	157	VVLYKRYCKFINGWLI----MSSMLLELFETYIYIGEVLKTYNNAMDYPILLTNWFG	210
: :	: :	: :	
Db	65	SV--AHTCDVNVTGWLRLHANGASSMFVICLHLGRGIYGSY-----LYKETWN 114	

DT	01-MAY-1999 (TREMBrel. 10, Last annotation update)	DE CYTOCHROME B LIGHT STRAND.
OS	Antechinus naso (Antechinus habbema).	OS Didelphis marsupialis.
OG	Mitochondrion.	OG Mitochondrion.
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;	OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC	Metatheria; Dasyuromorpha; Dasyuridae; Antechinus.	OC Metatheria; Dasyuromorpha; Dasyuridae; Antechinus.
RN	[1]	RN [1]
RP	SEQUENCE FROM N.A.	RP SEQUENCE FROM N.A.
RA	KRAJEWSKI C., PAINTER J., DRISKELL A.C., BUCKLEY L., WESTERMAN M.;	RA PATTON J.L., ROTS MARIA S.F., DA SILVA N.F.;
RL	Sci. New Guinea 19:157-166(1993).	RL J. Mammal. Evol. 3:3-29(1996).
RN	[2]	RL - ; CATALYTIC ACTIVITY: QH(2) + 2 FERRICYTOCHROME C = Q + 2
RP	SEQUENCE FROM N.A.	CC FERROCYTOCHROME C.
RA	KRAJEWSKI C., PAINTER J., BUCKLEY L., WESTERMAN M.;	CC - ; COFACTOR: TWO HEME GROUPS (B562 AND B566) WHICH ARE NOT COVALENTLY
RL	J. Mammal. Evol. 2:25-35(1996).	CC BOUND TO THE PROTEIN (BY SIMILARITY).
RN	[3]	DR EMBL: U34665; AAA99746.1; -
RP	SEQUENCE OF 1-287 FROM N.A.	DR PFAM; PF0032; cytochrome_b_C; 1.
RA	KRAJEWSKI C., BUCKLEY L., WOOLLEY P.A., WESTERMAN M.;	DR PFAM; PF0033; cytochrome_b_N; 1.
RL	J. Mammal. Evol. 3:81-91(1996).	KW Mitochondrion; Electron transport; Respiratory chain; Transmembrane; Hemee.
RN	[4]	SQ SEQUENCE 382 AA; 43139 MW; 907FBCA3 CRC32;
RA	SEQUENCE FROM N.A.	Query Match Score 4.6%; Length 382;
RA	ARMSTRONG L.A., KRAJEWSKI C., WESTERMAN M.;	Best Local Similarity 20.5%; Pred. No. 14;
RL	J. Mammal. 0:0-0(1998).	Matches 79; Conservative 53; Mismatches 115; Gaps 18;
RP	SEQUENCE FROM N.A.	QY 97 LCMTIVVATIKSVRYTEKNGQLIYPTEDEPSVQQLRNSVLNTLIMISIVVVMTFL 156
RA	KRAJEWSKI C.; Submitted (MAR-1994) to the EMBL/GenBank/DBJ databases.	Db 39 MCIIQIILT-----GLFLAMHYTSPT-----LTAFAS 64
RL	[6]	QY 157 VVLIKYRCYKPIFHGWLIM----SSLMLLEPLFTTYLGELYLKTYNVAMYDPTILLTVWN 210
DR	Submitted (FEB-1998) to the EMBL/GenBank/DBJ databases.	Db 65 SV--AHICRDVNYGWIRNIHANGASMFMGLFLHGRIYGSY-----LYKETWN 114
EMBL; U07576; AAC0634.1; -	PFAM; PF0032; cytochrome_b_C; 1.	QY 211 FGAVGMVCIHWKGPLYLQOAYLIMISALMAVFKYLPEWNSAWVILGAISYDLYAVLCP 270
DR	PFAM; PF0033; cytochrome_b_N; 1.	Db 115 IG-----VILLTVMATAFVGIVLPWQMSFGATVITNLSSAIPY 155
Mitochondrion.	Mitochondrion.	QY 271 KGPLRMLVE-----TAQERNEPFPALIYSAMWVTGMAK-----LDP 309
SEQUENCE 381 AA; 42738 MW; 9C6998AO CRC32;	Db 156 IG--NTLVENWGFSVDKATLTFPFLFILMMVYHLLFHETGSNNPTGLDP 213	
Query Match Score 4.6%; Length 381;	QY 310 SSQGALQPKD-----EMEEDSYDSFGEPSYPEVFEP--PLTGPFG-----	
Best Local Similarity 21.3%; Pred. No. 1.3;	Db 214 NSD--KIPHPHYTIKIDLGFLMILLSLAMESPDPLLSD---PDNFTPANPLNTPPH 267	
Matches 76; Conservative 55; Mismatches 111; Gaps 17;	QY 350 EEELEEEERGVKLGLDFIF-YSVLYGKAATGSGDWNTTLACFVATLIGCLTLILAV 408	
QY 138 SVLNNTLIMISIVVMTFLVVL-----KRYCYKFHOWLIM----SSLMLL 180	Db 268 IKP-----WYFLFAYLERSIPNKLGG----VILLASLILLIMPLHTST 311	
Db 35 SLIGACLIIQIL--MGLFLAMHITSDTLTAFTSSVAHICRDVNYGWIRNIHANGASMFM 92	QY 409 EKKALPALISITFGHLIFYFFTDNLV 434	
QY 181 FLFTYIYGEVLKYVNVAMDPTLLLTVNGFAGVMC1THWKGPLVQOAYLIMISALMA 240	Db 312 -QRSMFRPISQT--LFWMLTANLI 333	
Db 93 CLFHMGRCGIYGSY-----LYKETWNIG-----VILLTVWA 125	Search completed: March 18, 2000, 22:07:43	
QY 241 LYFIKYLPEWSAWVILGAISYDLYAVLCPKGPLRMLVE-----TAQERNEP 287	Job time: 123 sec	
Db 126 TAVFGYVLPWQMSFGATVITNLSSAIPYIG--TTLAZWVWGGFAVDKATLTFFAFH 183		
QY 288 IFPALIYSSAMWVTGMA-----AKLDPSSQCALQPYDPE-----323		
Db 184 ILPEIVVAAVILVHLETHETGSNNPSGLNPSD-----KLFPHPYTTIKDALGMNTLLAI 240		
QY 323 -MEEDSYDSFGEPSYPEVFEP--PLTGPFGEELEBEEERGVKLGLDFIF-YSVLYGKA 378		
Db 241 LLALFSPDSLGD---PDNESPANLNTPHKPE-----WYFLFAYLERSIPNKLN 286		
QY 379 ATGSGDWNTTLACFVATLIGCLTLILAVFKALPALISITFGHLIFYFFTDNLV 434		
Db 287 KLGG-----VLLASLILLIPLHHTA-NORSMMRFDISQT--LFWMLTANLI 333		
RESULT 15		
Q34340 PRELIMINARY; PRT; 382 AA.		
ID Q34340		
AC Q34340;		
DT 01-NOV-1996 (TREMBrel. 01, Created)		
DT 01-NOV-1996 (TREMBrel. 01, Last sequence update)		
DT 01-MAY-1999 (TREMBrel. 10, Last annotation update)		

=> e

E6 1 PRESENILIN-2 PS2S (MOUSE PS-2SHORT ISOFORM) /CN
E7 1 PRESENILIN-ASSOCIATED PROTEIN 1 (HPAP-1) (HUMAN INCYTE
CLONE 1353337) /CN
E8 1 PRESENILINASE/CN
E9 1 PRESEP-AGRI/CN
E10 1 PRESER ACE/CN
E11 1 PRESERCAR/CN
E12 1 PRESERIN M 72/CN
E13 1 PRESERIN T 72/CN
E14 1 PRESERT/CN
E15 1 PRESERVAC WETPROOF/CN
E16 1 PRESERVAL/CN
E17 1 PRESERVAL B/CN

=> s e4-6

1 "PRESENILIN-2 (HUMAN ISOFORM)"/CN
1 "PRESENILIN-2 PS2CCAS (MOUSE)"/CN
1 "PRESENILIN-2 PS2S (MOUSE PS-2SHORT ISOFORM)"/CN
L1 3 ("PRESENILIN-2 (HUMAN ISOFORM)"/CN OR "PRESENILIN-2 PS2CCAS
(MOUSE)"/CN OR "PRESENILIN-2 PS2S (MOUSE PS-2SHORT
ISOFORM)"/CN)

=> d 1-3 ide can

L1 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2000 ACS
RN 251358-30-2 REGISTRY
CN Presenilin-2 (human isoform) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN 5: PN: WO9960122 SEQID: 5 claimed protein
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
*** USE 'SQD' OR 'SQIDE' FORMATS TO DISPLAY SEQUENCE ***
1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 132:11416

L1 ANSWER 2 OF 3 REGISTRY COPYRIGHT 2000 ACS
RN 200445-64-3 REGISTRY
CN 330-448-presenilin-2 (Mus musculus isoform PS2Ccas) (9CI) (CA INDEX
NAME)
OTHER NAMES:
CN Presenilin-2 PS2Ccas (mouse)
FS PROTEIN SEQUENCE
MF Unspecified
CI MAN
SR CA
LC STN Files: CA, CAPLUS

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***